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L1 ~~FILE 'REGISTRY'~~ ENTERED AT 12:05:21 ON 30 AUG 2002
918 S [EG][AT][GV][WG][PS]S/SQSP

L2 ~~FILE 'HCAPLUS'~~ ENTERED AT 12:06:28 ON 30 AUG 2002
491 S L1
L3 7 S L2 AND MICROTI

L3 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2001:833517 HCAPLUS

DOCUMENT NUMBER: 135:367756

TITLE: Babesia **microti** antigens and methods
for the diagnosis and treatment of Babesia
microti infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton,
Raymond L.; Sleath, Paul R.; McNeill, Patricia
D.; Homer, Mary J.; Secrist, Heather

PATENT ASSIGNEE(S): Corixa Corporation, USA

SOURCE: PCT Int. Appl., 195 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 7

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001085947	A2	20011115	WO 2001-US15192	20010509
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
US 2001029295	A1	20011011	US 2000-737178	20001213
PRIORITY APPLN. INFO.:			US 2000-569098	A 20000510
			US 2000-605724	A 20000627
			US 2000-656688	A 20000907
			US 2000-685436	A 20001010
			US 2000-737178	A 20001213
			US 2001-794764	A 20010226
			US 1996-723142	A2 19961001
			US 1997-845258	A2 19970424
			US 1997-990571	A2 19971211
			WO 1998-US26437	A2 19981211
			US 1999-286488	A2 19990405
			US 2000-528784	A2 20000317
			WO 2000-US9136	A2 20000405
AB	Compds. and methods for the diagnosis and treatment of B. microti infection are disclosed. The compds. provided include polypeptides that contain at least one antigenic portion of a B. microti antigen and DNA sequences encoding such polypeptides. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compns. and immunogenic			

Claim 3
Sept 1-DU(135)
General
Gallen
for B
amb
2001/0509

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compsns. comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of *B. microti* infection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

- IT 206205-11-0, Antigen BMNI-1 (*Babesia microti*) *Peak*
 206205-12-1, Antigen BMNI-2 (*Babesia microti*)
 206205-13-2, Antigen BMNI-3 (*Babesia microti*)
 206205-16-5, Antigen BMNI-6 (*Babesia microti*)
 206205-20-1, Antigen BMNI-12 (*Babesia microti*)
 206205-21-2, Antigen BMNI-13 (*Babesia microti*)
 206205-23-4, Antigen BMNI-16 (*Babesia microti*)
 RL: BUU (Biological use, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (amino acid sequence; of *Babesia microti* antigens used in diagnosis and treatment of *Babesia microti* infection)
- IT 206205-33-6 206205-35-8 206205-36-9
 227296-22-2 227296-23-3 227296-26-6
 227296-30-2 227296-31-3 227296-32-4
 227296-33-5 227296-34-6 227296-35-7,
 Antigen MN2 (*Bombesia microtia* fragment) 227296-36-8
 227296-37-9, Antigen MN3 (*Bombesia microtia* fragment)
 RL: PRP (Properties)
 (unclaimed protein sequence; *babesia microti* antigens and methods for the diagnosis and treatment of *Babesia microti* infection)
- IT 205488-48-8 205488-54-6 334074-87-2
 334074-88-3 334074-89-4 334074-90-7
 RL: PRP (Properties)
 (unclaimed sequence; *babesia microti* antigens and methods for the diagnosis and treatment of *Babesia microti* infection)

L3 ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2001:748300 HCAPLUS

DOCUMENT NUMBER: 135:299589

TITLE: Nucleic acids and proteins for the diagnosis and treatment of *Babesia microti* infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton, Raymond L.; Sleath, Paul R.; McNeill, Patricia D.

PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 26 pp., Cont.-in-part of U. S. Ser. No. 685,436.
 CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 7

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2001029295	A1	20011011	US 2000-737178	20001213
US 6306396	B1	20011023	US 1996-723142	19961001
US 6183976	B1	20010206	US 1997-845258	19970424
US 6214971	B1	20010410	US 1997-990571	19971211

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WO 9929869 A1 19990617 WO 1998-US26437 19981211
W: AU, CA, JP, MX, NZ
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
NL, PT, SE
WO 2000060090 A1 20001012 WO 2000-US9136 20000405
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN,
CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,
US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF,
BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
WO 2001085947 A2 20011115 WO 2001-US15192 20010509
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,
GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR,
TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ,
MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH,
CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE,
TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD,
TG

PRIORITY APPLN. INFO.:

US 1996-723142 A2 19961001
US 1997-845258 A2 19970424
US 1997-990571 A2 19971211
WO 1998-US26437 A2 19981211
US 1999-286488 A2 19990405
US 2000-528784 A2 20000317
WO 2000-US9136 A2 20000405
US 2000-569098 A2 20000510
US 2000-605724 A2 20000627
US 2000-656688 A2 20000907
US 2000-685436 A2 20001010
US 2000-737178 A 20001213
US 2001-794764 A 20010226

AB Compds. and methods for the diagnosis and treatment of B.
microtiinfection are disclosed. The compds. provided include
polypeptides that contain at least one antigenic portion of a B.
microtiantigen and DNA sequences encoding such polypeptides.
Antigenic epitopes of such antigens are also provided, together with
pharmaceutical compns. and immunogenic compns. comprising such
polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits
contg. such polypeptides, DNA sequences or antigenic epitopes and a
suitable detection reagent may be used for the detection of B.
microtiinfection in patients and biol. samples. Antibodies directed
against such polypeptides and antigenic epitopes are also provided.

IT 206205-36-9

RL: BPR (Biological process); BSU (Biological study, unclassified);
PRP (Properties); BIOL (Biological study); PROC (Process)
(nucleic acids and proteins for the diagnosis and treatment of
Babesia microti infection)

IT 227296-26-6 227296-30-2 227296-31-3

227296-32-4 227296-34-6 227296-35-7,

Antigen MN2 (Bombesia microtia fragment) 227296-36-8

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227296-37-9, Antigen MN3 (Bombesia microtia fragment)
334074-87-2 334074-88-3 334074-89-4
334074-90-7

RL: PRP (Properties)

(unclaimed protein sequence; nucleic acids and proteins for the
diagnosis and treatment of Babesia **microti** infection)

L3 ANSWER 3 OF 7 HCAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2001:255942 HCAPLUS

DOCUMENT NUMBER: 134:294507

TITLE: Compounds and methods for the diagnosis and
treatment of Babesia **microti** infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton,
Raymond

PATENT ASSIGNEE(S): Corixa Corporation, USA

SOURCE: U.S., 78 pp., Cont.-in-part of U.S. Ser. No.
845,258.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 7

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6214971	B1	20010410	US 1997-990571	19971211
US 6306396	B1	20011023	US 1996-723142	19961001
US 6183976	B1	20010206	US 1997-845258	19970424
WO 9929869	A1	19990617	WO 1998-US26437	19981211
W: AU, CA, JP, MX, NZ				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
AU 9918204	A1	19990628	AU 1999-18204	19981211
US 2001029295	A1	20011011	US 2000-737178	20001213

PRIORITY APPLN. INFO.:

US 1996-723142	A2	19961001
US 1997-845258	A2	19970424
US 1997-990571	A	19971211
WO 1998-US26437	W	19981211
US 1999-286488	A2	19990405
US 2000-528784	A2	20000317
WO 2000-US9136	A2	20000405
US 2000-569098	A2	20000510
US 2000-605724	A2	20000627
US 2000-656688	A2	20000907
US 2000-685436	A2	20001010

AB Compds. and methods for the diagnosis and treatment of Babesia
microti infection are disclosed. The compds. provided
include polypeptides that contain at least one antigenic portion of
a B. **microti** antigen and DNA sequences encoding such
polypeptides. Antigenic epitopes of such antigens are also
provided, together with pharmaceutical compns. and vaccines
comprising such polypeptides, DNA sequences or antigenic epitopes.
Diagnostic kits contg. such polypeptides, DNA sequences or antigenic
epitopes and a suitable detection reagent may be used for the
detection of B. **microti** infection in patients and biol.
samples. Antibodies directed against such polypeptides and
antigenic epitopes are also provided.

IT 206205-11-0 206205-12-1 206205-13-2

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206205-16-5 206205-20-1 206205-21-2

206205-23-4

RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(amino acid sequence; compds. and methods for the diagnosis and treatment of Babesia **microti** infection)

IT 206205-33-6 206205-35-8 206205-36-9

227296-22-2 227296-23-3 227296-26-6

227296-30-2 227296-31-3 227296-32-4

227296-33-5 227296-34-6 227296-35-7,

Antigen MN2 (Babesia microtia fragment) 227296-36-8

227296-37-9, Antigen MN3 (Babesia microtia fragment)

334074-87-2

RL: PRP (Properties)

(unclaimed protein sequence; compds. and methods for the diagnosis and treatment of Babesia **microti** infection)

IT 205488-48-8 205488-54-6 334074-88-3

334074-89-4 334074-90-7

RL: PRP (Properties)

(unclaimed sequence; compds. and methods for the diagnosis and treatment of Babesia **microti** infection)

REFERENCE COUNT: 13 THERE ARE 13 CITED REFERENCES AVAILABLE
FOR THIS RECORD. ALL CITATIONS AVAILABLE
IN THE RE FORMAT

L3 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2001:91448 HCAPLUS

DOCUMENT NUMBER: 134:158493

TITLE: Nucleic acids and proteins for the diagnosis and treatment of Babesia **microti** infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton, Raymond; Sleath, Paul R.

PATENT ASSIGNEE(S): Corixa Corporation, USA

SOURCE: U.S., 66 pp., Cont.-in-part of U.S. Ser. No. 723,142.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 7

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6183976	B1	20010206	US 1997-845258	19970424
US 6306396	B1	20011023	US 1996-723142	19961001
EP 834567	A2	19980408	EP 1997-117067	19971001
EP 834567	A3	19990331		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI

US 6214971 B1 20010410 US 1997-990571 19971211

US 2001029295 A1 20011011 US 2000-737178 20001213

PRIORITY APPLN. INFO.: US 1996-723142 A2 19961001

US 1997-845258 A 19970424

US 1997-990571 A2 19971211

WO 1998-US26437 A2 19981211

US 1999-286488 A2 19990405

US 2000-528784 A2 20000317

WO 2000-US9136 A2 20000405

Searcher : Shears 308-4994

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US 2000-569098 A2 20000510
US 2000-605724 A2 20000627
US 2000-656688 A2 20000907
US 2000-685436 A2 20001010

OTHER SOURCE(S): MARPAT 134:158493

AB Compds. and methods for the diagnosis and treatment of Babesia **microti** infection are disclosed. The compds. provided include polypeptides that contain at least one antigenic portion of a B. **microti** antigen and DNA sequences encoding such polypeptides. The DNA sequences encoding B. **microti** antigens were prepd. by screening a B. **microti** expression library with sera obtained from patients infected with B. **microti**. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compns. and vaccines comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of B. **microti** infection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

IT 205488-48-8 205488-54-6 206205-11-0
206205-12-1 206205-13-2 206205-16-5
206205-20-1 206205-21-2 206205-23-4
206205-33-6 206205-35-8 206205-36-9

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); USES (Uses)

(amino acid sequence; nucleic acids and proteins for the diagnosis and treatment of Babesia **microti** infection)

REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2000:282570 HCAPLUS

DOCUMENT NUMBER: 133:72606

TITLE: Serological expression cloning of novel immunoreactive antigens of Babesia **microti**

AUTHOR(S): Lodes, Michael J.; Houghton, Raymond L.; Bruinsma, Elizabeth S.; Mohamath, Raodoh; Reynolds, Lisa D.; Benson, Darin R.; Krause, Peter J.; Reed, Steven G.; Persing, David H.

CORPORATE SOURCE: Corixa Corporation, Seattle, WA, 98104, USA
SOURCE: Infection and Immunity (2000), 68(5), 2783-2790
CODEN: INFIBR; ISSN: 0019-9567

PUBLISHER: American Society for Microbiology

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Increased recognition of the prevalence of human babesiosis in the United States, together with rising concern about the potential for transmission of this infection by blood transfusion, has provided motivation to develop definitive serol. and mol. tests for the causative agent, Babesia **microti**. To develop more sensitive and specific assays for B. **microti**, the authors screened a genomic expression library with patient serum pools. This screening resulted in the identification of three classes of novel genes and an addnl. two novel, unrelated genes, which together

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encode a total of 17 unique *B. microti* antigens. The first class (BMN1-2 family) of genes encodes seven closely related antigens with a degenerate six-amino-acid repeat that shows limited homol. to *Plasmodium* sp. merozoite and sporozoite surface antigens. A second class (BMN1-8 family) of genes encodes six related antigens, and the third class (BMN1-17 family) of genes encodes two related antigens. The two remaining genes code for novel and unrelated sequences. Among the three classes of antigens and remaining novel sequences, five were chosen to code for the most immunodominant antigens (BMN1-2, -9, -15, and -17 and MN-10). Western blot anal. with the resulting recombinant proteins indicated that these antigens were targets of humoral immune responses during *B. microti* infection in humans.

IT 278626-95-2 278626-96-3 278626-97-4
278626-98-5 278626-99-6

RL: BPR (Biological process); BSU (Biological study, unclassified);
PRP (Properties); BIOL (Biological study); PROC (Process)
(amino acid sequence; serol. expression cloning of novel
immunoreactive antigens of *Babesia microti*)

REFERENCE COUNT: 36 THERE ARE 36 CITED REFERENCES AVAILABLE
FOR THIS RECORD. ALL CITATIONS AVAILABLE
IN THE RE FORMAT

L3 ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1999:388301 HCAPLUS

DOCUMENT NUMBER: 131:85409

TITLE: Antigen and gene sequences for diagnosis and
treatment of *Babesia microti* infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton,
Raymond; Sleath, Paul R.; Persing, David;
Bruinsma, Elizabeth

PATENT ASSIGNEE(S): Corixa Corporation, USA; Mayo Foundation for
Medical Education and Research

SOURCE: PCT Int. Appl., 126 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 7

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9929869	A1	19990617	WO 1998-US26437	19981211
W: AU, CA, JP, MX, NZ				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 6214971	B1	20010410	US 1997-990571	19971211
AU 9918204	A1	19990628	AU 1999-18204	19981211
US 2001029295	A1	20011011	US 2000-737178	20001213
PRIORITY APPLN. INFO.:			US 1997-990571	A 19971211
			US 1996-723142	A2 19961001
			US 1997-845258	A2 19970424
			WO 1998-US26437	W 19981211
			US 1999-286488	A2 19990405
			US 2000-528784	A2 20000317
			WO 2000-US9136	A2 20000405
			US 2000-569098	A2 20000510
			US 2000-605724	A2 20000627

Searcher : Shears 308-4994

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US 2000-656688 A2 20000907

US 2000-685436 A2 20001010

AB Compds. and methods for the diagnosis and treatment of Babesia **microti** infection are disclosed. The compds. provided include polypeptides that contain at least one antigenic portion of a B. **microti** antigen and DNA sequences encoding such polypeptides. Nine antigens share some homol., contain a degenerate repeat of six amino acids with 9-22 repeats occurring in each antigen, and bear some similarity to a Plasmodium falciparum merozoite surface antigen gene. A second group of five antigens bear some homol. to each other but do not show homol. to any previously identified sequences. Two synthetic peptides (BABS-1 and BABS-4) were made to the repeat region of isolated B. **microti** antigen BMNI-3. Twelve BMNI-6 homologs were obtained from hamster and human patients. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compns. and vaccines comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of B. **microti** infection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

IT 205488-48-8

RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(BABS-1 peptide fragment of repeat region of BMNI-3; antigen and gene sequences for diagnosis and treatment of Babesia **microti** infection)

IT 205488-54-6

RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(BABS-4 peptide fragment of repeat region of BMNI-3; antigen and gene sequences for diagnosis and treatment of Babesia **microti** infection)

IT 206205-11-0 206205-12-1 206205-13-2

206205-16-5 206205-20-1 206205-21-2

206205-33-6 206205-35-8 227296-22-2

227296-23-3 227296-26-6 227296-30-2

227296-31-3 227296-32-4 227296-33-5

227296-34-6 227296-35-7, Antigen MN2 (Bombesia

microtia fragment) 227296-36-8 227296-37-9,

Antigen MN3 (Bombesia microtia fragment)

RL: ANT (Analyte); BOC (Biological occurrence); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); OCCU (Occurrence); USES (Uses)

(amino acid sequence; antigen and gene sequences for diagnosis and treatment of Babesia **microti** infection)

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1998:229027 HCAPLUS

DOCUMENT NUMBER: 128:292989

TITLE: Antigens of Babesia microtia and their use in the diagnosis and treatment of infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton,

Searcher : Shears 308-4994

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09/853079

PATENT ASSIGNEE(S): Raymond; Sleath, Paul R.
SOURCE: Corixa Corp., USA
Eur. Pat. Appl., 113 pp.
CODEN: EPXXDW
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 7
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 834567	A2	19980408	EP 1997-117067	19971001
EP 834567	A3	19990331		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
US 6306396	B1	20011023	US 1996-723142	19961001
US 6183976	B1	20010206	US 1997-845258	19970424
PRIORITY APPLN. INFO.:			US 1996-723142	A 19961001
			US 1997-845258	A 19970424

AB Antigens and epitopes of *Babesia microti* that can be used in the diagnosis and treatment of infection are described. Genes for the antigens or antibodies against them can be used in the detection of *B. microti*. CDNAS for these antigens were cloned by screening an expression library in .lambda.ZAP with antiserum to *B. microti*.

IT 205488-48-8 205488-54-6 206205-11-0
206205-12-1 206205-13-2 206205-16-5
206205-20-1 206205-21-2 206205-23-4
206205-33-6 206205-35-8 206205-36-9
RL: ANT (Analyte); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
(amino acid sequence; antigens of *Babesia microti* and their use in diagnosis and treatment of infection)

E1 THROUGH E32 ASSIGNED

~~FILE 'REGISTRY'~~ ENTERED AT 12:07:45 ON 30 AUG 2002

L4 32 SEA FILE=REGISTRY ABB=ON PLU=ON (205488-48-8/BI OR
205488-54-6/BI OR 206205-11-0/BI OR 206205-12-1/BI OR
206205-13-2/BI OR 206205-16-5/BI OR 206205-20-1/BI OR
206205-21-2/BI OR 206205-33-6/BI OR 206205-35-8/BI OR
206205-36-9/BI OR 206205-23-4/BI OR 227296-26-6/BI OR
227296-30-2/BI OR 227296-31-3/BI OR 227296-32-4/BI OR
227296-34-6/BI OR 227296-35-7/BI OR 227296-36-8/BI OR
227296-37-9/BI OR 227296-22-2/BI OR 227296-23-3/BI OR
227296-33-5/BI OR 334074-87-2/BI OR 334074-88-3/BI OR
334074-89-4/BI OR 334074-90-7/BI OR 278626-95-2/BI OR
278626-96-3/BI OR 278626-97-4/BI OR 278626-98-5/BI OR
278626-99-6/BI)

L5 32 L4 AND L1

L5 ANSWER 1 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 334074-90-7 REGISTRY
CN 90: PN: US6214971 FIGURE: 6 unclaimed sequence (9CI) (CA INDEX
NAME)
OTHER NAMES:

Searcher : Shears 308-4994

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09/853079

CN 10: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CN 204: PN: WO0185947 FIGURE: 6 unclaimed sequence
CI MAN
SQL 147

```
SEQ      1 YITLFLMSGV VFAGDTDREA GGPSGTVGPS EAGGPSEAGG PSEAGGPSEA
          == =====
          51 GGPSEAGGPS EAGGPSHAGG PSEAGGPSGT GWPSEAGWPS EAGWPSEAGW
             =====
          101 PSEAGWPSEA GWPSERFQYQ LLWYSRRIVI FNEIYLSHIY EHSVMIL
             =====
```

HITS AT: 19-66, 73-114

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

L5 ANSWER 2 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN **334074-89-4** REGISTRY
CN 86: PN: US6214971 FIGURE: 6 unclaimed sequence (9CI) (CA INDEX
NAME)

OTHER NAMES:

CN 200: PN: WO0185947 FIGURE: 6 unclaimed sequence
CN 6: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CI MAN
SQL 138

```
SEQ      1 AGDTDREAGG PSGTVGPSSA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
          =====
          51 GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA GGPSEAGWPS EAGWPSEAGG
             =====
          101 PSGTGWPSEA GWPSEAGWPS EAGWPSEAGW PSERFGYQ
             =====
```

HITS AT: 7-18, 25-132

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

L5 ANSWER 3 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN **334074-88-3** REGISTRY
CN 81: PN: US6214971 FIGURE: 6 unclaimed sequence (9CI) (CA INDEX
NAME)

OTHER NAMES:

CN 195: PN: WO0185947 FIGURE: 6 unclaimed sequence
CN 1: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CI MAN
SQL 111

```
SEQ      1 GDTDREAGGP SGTVPSEAG GPSEAGGPSG TVGPSEAGGP SEAGGPSGTG
          =====
          51 WPSEAGGPSG TVGPSEAGGP SEAGGPSGTG WPSGTGWPSE VGWPSERFGY
             =====
          101 QLLWYSRRIV I
```

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09/853079

HITS AT: 6-89

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

L5 ANSWER 4 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN **334074-87-2** REGISTRY

CN 80: PN: US6214971 FIGURE: 6 unclaimed protein (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 194: PN: WO0185947 FIGURE: 6 unclaimed sequence

CN 1: PN: US20010029295 FIGURE: 6A-6B unclaimed protein

CI MAN

SQL 112

SEQ 1 AGDTDREAGG PSGTVGPSEA GGPSEAGGPS GTVGPSEAGG PSEAGGPSGT

51 GWPSEAGGPS GTVGPSEAGG PSEAGGPSGT GWPSGTGWPS EVGWPSERFG

101 YQLLWYSRRI VI

HITS AT: 7-90

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

L5 ANSWER 5 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN **278626-99-6** REGISTRY

CN Antigen BMN1-13 (Babesia microtia strain MN1 clone bmn1-13 precursor) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN GenBank AF206524-derived protein GI 7716011

CI MAN

SQL 262

SEQ 1 MVSFKSILVP YITLFLMSGV VFASDTDPEA GGPSGTVGPS EAGGPSEAGG

51 PSGTGWPSSEA GGPSEAGGPS GTGWPSEAGW SSERFGYQLL PYSRRIVTFN

101 EVCLSYIYKH SVMILERDRV NDGHKDYIEE KTKEKNKLKK ELEKCFPEQY

151 SLMKKEELAR IFDNASTISS KYKLLVDEIS NKAYGTLEGP AADNFDHFRN

201 IWKSIVLKDM FIYCDLLQH LIYKFYYDNT INDIKKNFDE SKSKALVLRD

251 KITKKDVYVN DH

HITS AT: 29-82

REFERENCE 1: 133:72606

L5 ANSWER 6 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN **278626-98-5** REGISTRY

CN Antigen BMN1-7 (Babesia microtia strain MN1 clone bmn1-7 precursor) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN GenBank AF206250-derived protein GI 7715998

CI MAN

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09/853079

SQL 289

```
SEQ      1 MVSFKSILVP YITLFLMSGV VFAGDTDREA GGPSGTVGPS EAGGPSEAGG
          == =====
        51 PSEAGGPSEA GGPSEAGGPS EAGGPSEAGG PSGTVGPSEA GGPSEAGGPS
          =====
       101 EAGGPSEAGW PSEAGWPSEA GWPSEAGWPS EAGWPSEAGW PSERFGYQLL
          =====
       151 WYSRRIVIFN EIYLSHIYEH SVMILERDRV NDGHKDYIEE KTKENKLNKK
       201 ELEKCFPEQY SLMKKEELAR IIDNASTISS KYKLLVDEIS NKAYGTLEGP
       251 AADDFDHFNR IWKSIVPKNM FLYCDLLLKH LIRLTPRKS
```

HITS AT: 29-142

REFERENCE 1: 133:72606

L5 ANSWER 7 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 278626-97-4 REGISTRY
CN Antigen BMN1-6 (Babesia microtia strain MN1 clone bmn1-6 precursor)
(9CI) (CA INDEX NAME)

OTHER NAMES:

CN GenBank AF206249-derived protein GI 7715996

CI MAN

SQL 298

```
SEQ      1 MVSFKSILVP YITLFLMSGV VFAGDTDREA GGPSGTVGPS EAGGPSEAGG
          == =====
        51 PSEAGGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSGT GWPSEAGWPS
          =====
       101 EAGWPSEAGW PSEAGWPSEA GWPSEAFGYQ LLWYSRRIVI FNEIYLSHIY
          =====
       151 EHSVMILERD RVNDGHKDYI EEKTKENKLN KKELEKCFPE QYSLMKKEEL
       201 ARIIDNASTI SSKYKLLVDE ISNKAYGTLE GPAADDFDHF RNIWKSIVPK
       251 NMFLYCDLLL KHLIRKFYCD NTINDIKKNF DDIEKLGCFQ ARSFLPVN
```

HITS AT: 29-124

REFERENCE 1: 133:72606

L5 ANSWER 8 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 278626-96-3 REGISTRY
CN Antigen BMN1-3 (Babesia microtia strain MN1 clone bmn1-3 precursor)
(9CI) (CA INDEX NAME)

OTHER NAMES:

CN GenBank AF206245-derived protein GI 7715986

CI MAN

SQL 362

```
SEQ      1 MVSFKSILVP YITLFLMSGV VFASDTDPEA GGPSEAGGPS GTVGPSEAGG
          == =====
        51 PSEAGGPSGT GWPSEAGGPS EAGGPSEAGG PSEAGGPSGT GWPSGTGWPS
          =====
       101 EAGWSSERFG YQLLPYSRRI VIFNEVCLSY IYKHSVMILE RDRVNDGHKD
          =====
       151 YIEEKTKEKN KKKKELEKCF PEQYSLMKKE ELARIFDNAS TISSKYKLLV
       201 DEISNKAYGT LEGPAADNFD HFRNIWKSIV LKDMFIYCDL LLQHLIYKFY
       251 YDNTVNDIKK NFDESKSKAL VLRDKITKGD GDYNTHFEDM IKELNSAAEE
       301 FNKIVDIMIS NIGDYDEYDS IASFKPFLSM ITEITKITKV SNVIIPGIKA
       351 LTLTVFLIFI TK
```

HITS AT: 29-106

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09/853079

REFERENCE 1: 133:72606

L5 ANSWER 9 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN **278626-95-2** REGISTRY
CN Antigen BMN1-2 (Babesia microtia strain MN1 clone bmn1-2 precursor)
(9CI) (CA INDEX NAME)
OTHER NAMES:
CN GenBank AF206244-derived protein GI 7715984
CI MAN
SQL 326

```
SEQ      1 MVSFKSILVP YITLFLMSGV VFASDTPDPEA GGPSEAGGPS GTVGPSEAGG
          == =====
        51 PSEAGGPSGT VGPSEAGGPS EAGGPSGTGW PSEAGGPSEA GGPSGTVGPS
          =====
       101 EAGGPSEAGG PSGTGWPSEA GGPSEAGGPS EAGGPSEAGG PSGTGWPSGT
          =====
       151 GWPSEAGWSS ERFYQLLPY SRRIVIFNEV CLSYIYKHSV MILERDRVND
          =====
       201 GHKDYIEEKT KEKNKLKKEL EKFPEQYSL MKKEELARIF DNASTISSKY
       251 KLLVDEISNK AYTLEGPAE DNFDHFRNIW KSIVLKDMFI YCDLLLQHLI
       301 YKFYYDNTVN DIKKNFDESW TQTLKE
```

HITS AT: 29-160

REFERENCE 1: 133:72606

L5 ANSWER 10 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN **227296-37-9** REGISTRY
CN Antigen MN3 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 11: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CN 47: PN: WO0185947 SEQID: 78 unclaimed protein
CN 74: PN: WO0060090 SEQID: 78 unclaimed protein
CN 78: PN: US6214971 SEQID: 78 unclaimed protein
CN 79: PN: US6214971 SEQID: 79 unclaimed protein
CN 91: PN: US6214971 FIGURE: 6 unclaimed sequence
CN 92: PN: US6214971 FIGURE: 6 unclaimed sequence
CN Antigen MRT (Bombesia microtia fragment)
CI MAN
SQL 120

```
SEQ      1 AGDTPREAGG PSGTGVPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
          =====
        51 GGPSEAGGPS EAGGPSEAGG PSGTGWPSEA GWPSEAGWPS EAGWPSEAGW
          =====
       101 PSEAGWPSEF FGYQLLWYSR
          =====
```

HITS AT: 7-108

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

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REFERENCE 5: 131:85409

L5 ANSWER 11 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-36-8 REGISTRY
CN Antigen MN1PAT (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 46: PN: WO0185947 SEQID: 77 unclaimed protein
CN 73: PN: WO0060090 SEQID: 77 unclaimed protein
CN 77: PN: US6214971 SEQID: 77 unclaimed protein
CN 89: PN: US6214971 FIGURE: 6 unclaimed sequence
CN 9: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CI MAN
SQL 113

SEQ 1 AGDTDREAGG PSGTVGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
=====

51	GGPSEAGGPS	EAGGPSGTGW	PSEAGWPSEA	GWPSEAGWPS	EAGWPSEAGW
	=====	=====	=====	=====	=====

101 PSERFGYQLL WYS
==

HITS AT: 7-102

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

L5 ANSWER 12 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-35-7 REGISTRY
CN Antigen MN2 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 45: PN: WO0185947 SEQID: 76 unclaimed protein
CN 72: PN: WO0060090 SEQID: 76 unclaimed protein
CN 76: PN: US6214971 SEQID: 76 unclaimed protein
CN 88: PN: US6214971 FIGURE: 6 unclaimed sequence
CN 8: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CI MAN
SQL 94

SEQ 1 AGDTDREAGG PSGTVGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
=====

51	GGPSEAGGPS	GTGWPSEAGW	PSEAGWPSEA	GWPSEAGWPS	EAGW
	=====	=====	=====	=====	=====

HITS AT: 7-90

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

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09/853079

REFERENCE 5: 131:85409

L5 ANSWER 13 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-34-6 REGISTRY
CN Antigen MN1HAM (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 44: PN: WO0185947 SEQID: 75 unclaimed protein
CN 71: PN: WO0060090 SEQID: 75 unclaimed protein
CN 75: PN: US6214971 SEQID: 75 unclaimed protein
CN 7: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CN 87: PN: US6214971 FIGURE: 6 unclaimed sequence
CI MAN
SQL 118

```
SEQ      1 AGDTDREAGG PSGTVGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
          =====
        51 GGPSEAGGPS EAGGPSGTGW PSEAGWPSEA GWPSEAGWPS EAGWPSEAGW
          =====
       101 PSERFGYQLL WYSRRIVI
          ==
```

HITS AT: 7-102

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

L5 ANSWER 14 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-33-5 REGISTRY
CN Antigen RIFS (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 43: PN: WO0185947 SEQID: 74 unclaimed protein
CN 70: PN: WO0060090 SEQID: 74 unclaimed protein
CN 74: PN: US6214971 SEQID: 74 unclaimed protein
CI MAN
SQL 138

```
SEQ      1 AGDTDREAGG PSGTVGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
          =====
        51 GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA GGPSEAGWPS EAGWPSEAGG
          =====
       101 PSGTGWPSEA GWPSEAGWPS EAGWPSEAGW PSERFGYQ
          =====
```

HITS AT: 7-132

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

REFERENCE 3: 133:295360

REFERENCE 4: 131:85409

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09/853079

L5 ANSWER 15 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-32-4 REGISTRY
CN Antigen BI2018 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 42: PN: WO0185947 SEQID: 73 unclaimed protein
CN 5: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CN 69: PN: WO0060090 SEQID: 73 unclaimed protein
CN 73: PN: US6214971 SEQID: 73 unclaimed protein
CN 85: PN: US6214971 FIGURE: 6 unclaimed sequence
CI MAN
SQL 136

SEQ 1 GDTDREAGGP SGTVGPSEAG GPSEAGGPSE AGGPSEAGGP SEAGGPSEAG
=====

51 GPSEAGGPSE AGGPSEAGGP SEAGGPSEAG WPSEAGWPSE AGGPGSGTGWP
=====

101 SEAGWPSEAG WPSEAGWPSE AGWPSEFSGY QLLWYS
=====

HITS AT: 6-125

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

L5 ANSWER 16 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-31-3 REGISTRY
CN Antigen BI2253 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 41: PN: WO0185947 SEQID: 72 unclaimed protein
CN 4: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CN 68: PN: WO0060090 SEQID: 72 unclaimed protein
CN 72: PN: US6214971 SEQID: 72 unclaimed protein
CN 84: PN: US6214971 FIGURE: 6 unclaimed sequence
CI MAN
SQL 116

SEQ 1 EAGGPSTGTVG PSEAGGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
=====

51 GGPSEAGGPS EAGGPSEAGW PSEAGWPSEA GGPGSGTGWPS EAGWPSEAGW
=====

101 PSEAGWPSEA GWPSE
=====

HITS AT: 1-114

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

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09/853079

REFERENCE 5: 131:85409

L5 ANSWER 17 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-30-2 REGISTRY
CN Antigen BI2259 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 3: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CN 40: PN: WO0185947 SEQID: 71 unclaimed protein
CN 71: PN: US6214971 SEQID: 71 unclaimed protein
CN 83: PN: US6214971 FIGURE: 6 unclaimed sequence
CI MAN
SQL 136

```
SEQ      1 AGDTDREAGG PSGTVGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
          =====
        51 GGPSEAGGPS EAGGPSEAGG PSEAGWPSEA GWPSEAGGPS GTGWPSEAGW
          =====
       101 PSEAGWPSEA GWPSEAGWPS ERFQYQLLWY SRRIVI
          =====
```

HITS AT: 7-120

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

L5 ANSWER 18 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-26-6 REGISTRY
CN Antigen BI2227 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 2: PN: US20010029295 FIGURE: 6A-6B unclaimed protein
CN 39: PN: WO0185947 SEQID: 70 unclaimed protein
CN 67: PN: WO0060090 SEQID: 70 unclaimed protein
CN 70: PN: US6214971 SEQID: 70 unclaimed protein
CN 82: PN: US6214971 FIGURE: 6 unclaimed sequence
CI MAN
SQL 118

```
SEQ      1 AGDTDREAGG PSGTVGPSEA GGPSEAGGPS EAGGPSEAGG PSEAGGPSEA
          =====
        51 GGPSEAGGPS EAGGPSEAGG PSEAGWPSEA GWPSEAGGPS GTGWPSEAGW
          =====
       101 PSEAGWPSEA GWPSEAGW
          =====
```

HITS AT: 7-114

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 133:295360

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09/853079

REFERENCE 5: 131:85409

L5 ANSWER 19 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-23-3 REGISTRY
CN Antigen BI1053 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 38: PN: WO0185947 SEQID: 69 unclaimed protein
CN 66: PN: WO0060090 SEQID: 69 unclaimed protein
CN 69: PN: US6214971 SEQID: 69 unclaimed protein
CI MAN
SQL 105

SEQ 1 AGDTRDREAGG PSGTVGPSEA GGPSEAGGPS GTVGPSEAGG PSEAGGPSGT
=====

51	GWPSEAGGPS	GTVGPSEAGG	PSEAGGPSGT	GWPSGTGWPS	EVGWPNEPFG
	=====	=====	=====	=====	

101 YHLLW
HITS AT: 7-90

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

REFERENCE 3: 133:295360

REFERENCE 4: 131:85409

L5 ANSWER 20 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 227296-22-2 REGISTRY
CN Antigen BI254 (Bombesia microtia fragment) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 37: PN: WO0185947 SEQID: 68 unclaimed protein
CN 65: PN: WO0060090 SEQID: 68 unclaimed protein
CN 68: PN: US6214971 SEQID: 68 unclaimed protein
CI MAN
SQL 101

SEQ 1 AGDTRDREAGG PSGTVGPSEA GGPSEAGGPS GTVGPSEAGG PSEAGGPSGT
=====

51	GWPSEAGGPS	GTVGPSEAGG	PSEAGGPSGT	GWPSGTGWPS	EVGWPIEPFG
	=====	=====	=====	=====	

101 Y
HITS AT: 7-90

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

REFERENCE 3: 133:295360

REFERENCE 4: 131:85409

L5 ANSWER 21 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 206205-36-9 REGISTRY
CN Antigen (Babesia microtia 367-amino acid) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 18: PN: WO0185947 SEQID: 49 unclaimed protein

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09/853079

CN 2: PN: US20010029295 SEQID: 49 claimed protein
CN 46: PN: WO0060090 SEQID: 49 unclaimed protein
CN 49: PN: US6183976 SEQID: 49 claimed protein
CN 49: PN: US6214971 SEQID: 49 unclaimed protein
CN Antigen BMNI-3 (Babesia microtia isoform 2 fragment)
CI MAN
SQL 367

SEQ 1 MVSFKSILVP YITLFLMSGV VFASDTDPEA GGPSEAGGPS GTVGPSEAGG
=====

51 PSEAGGPSGT GWPSEAGGPS EAGGPSEAGG PSEAGGPSGT GSEAGGWPSG
=====

101 TGWPSEAGWS SERFGYQLLP YSRRIVIFNE VCLSYYIKHS VMILERDRVN
=====

151 DGHKDYIEEK TKEKNKLKKE LEKCFPEQYS LMKKEELARI FDNASTISSK
201 YKLLVDEISN KAYGTLEGPA ADNFDHFRNI WKSIVLKDMF IYCDLLLQHL
251 IYKFYYDNTV NDIKKNFDES KSKALVLRDK ITKKGDDYNT HFEDMIKELN
301 SAAEEFNKIV DIMISNIGDY DEYDSIASFK PFLSMITEIT KITKVSNNVII
351 PGIKALTTLTV FLIFITK

HITS AT: 29-88, 100-111

REFERENCE 1: 135:367756

REFERENCE 2: 135:299589

REFERENCE 3: 134:294507

REFERENCE 4: 134:158493

REFERENCE 5: 133:295360

REFERENCE 6: 128:292989

L5 ANSWER 22 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 206205-35-8 REGISTRY
CN Antigen (Babesia microtia 294-amino acid) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 14: PN: WO0185947 SEQID: 46 unclaimed protein
CN 45: PN: WO0060090 SEQID: 46 unclaimed protein
CN 46: PN: US6183976 SEQID: 46 claimed protein
CN 46: PN: US6214971 SEQID: 46 unclaimed protein
CN Antigen BMNI-7 (Babesia microtia isoform fragment)
CN Antigen BMNI-7 (Bomnesia microtia reverse complement fragment)
CI MAN
SQL 294

SEQ 1 LWFIKMVSFK SILVPYITLF LMSGAVFAGD TDREAGGPSG TVGPSEAGGP
=====

51 SEAGGPSEAG GPSEAGGPSE AGGPSEAGGP SEAGGPSGTV GPSEAGGPSE
=====

101 AGGPSEAGGP SEAGWPSEAG WPSEAGWPSE AGWPSEAGWP SEAGWPSERF
=====

151 GYQLLWYSRR IVIFNEIYLS HIYEHSMIL ERDRVNDGHK DYIEEKTKEK
201 NKLKKELEKC FPEQYSLMKK EELARIIDNA STISSKYKLL VDEISNKAYG
251 TLEGPAADDF DHFRNIWКСI VPKNNFLYCD LLLKHLIRLT PRKS

HITS AT: 34-147

REFERENCE 1: 135:367756

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09/853079

REFERENCE 2: 134:294507
REFERENCE 3: 134:158493
REFERENCE 4: 133:295360
REFERENCE 5: 131:85409
REFERENCE 6: 128:292989

L5 ANSWER 23 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 206205-33-6 REGISTRY
CN Antigen (Babesia microtia 154-amino acid) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 12: PN: WO0185947 SEQID: 44 unclaimed protein
CN 43: PN: WO0060090 SEQID: 44 unclaimed protein
CN 44: PN: US6183976 SEQID: 44 claimed protein
CN 44: PN: US6214971 SEQID: 44 unclaimed protein
CN Antigen BMNI-5 (Babesia microtia isoform 2 fragment)
CN Antigen BMNI-5 (Babesia microtia reverse complement 154-amino acid fragment)
CI MAN
SQL 154

SEQ 1 LWFIKMVSFK SILVPYITLF LMSGAVFAGD TDREAGGPSG TVGPSEAGGP
===== =====
51 SEAGGPSEAG GPSEAGGPSE AGGPSEAGGP SEAGGPSEAG GPSGTGWPSE
===== =====
101 AGGPSEAGGP SEAGGPSGTG WPSEAGWPSE AGWPSEAGWP SEAGWPSEAG
===== =====
151 WPSE
===

HITS AT: 34-153

REFERENCE 1: 135:367756
REFERENCE 2: 134:294507
REFERENCE 3: 134:158493
REFERENCE 4: 133:295360
REFERENCE 5: 131:85409
REFERENCE 6: 128:292989

L5 ANSWER 24 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 206205-23-4 REGISTRY
CN Antigen (Babesia microtia 128-amino acid) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 250: PN: WO0185947 SEQID: 31 claimed protein
CN 30: PN: WO0060090 SEQID: 31 unclaimed protein
CN 31: PN: US6183976 SEQID: 31 claimed protein
CN 31: PN: US6214971 SEQID: 31 claimed protein
CN Antigen BMNI-16 (Babesia microti)
CN Antigen BMNI-16 (Babesia microtia fragment)
CI MAN

Searcher : Shears 308-4994

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09/853079

SQL 128

```
SEQ      1 LWFIKMVSFK SILVPYITLF LMSGAVFASD TDPEAGGPSE AGGPSGTGVP
          =====
          51 SEAGGPSEAG GPSGTGWPSE AGGPSEAGGP SEAGGPSEAG GPSGTGWPSG
          =====
          101 TGWPSEAGWS SERFGYQLLP YSRRIVIF
          =====
```

HITS AT: 34-111

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

REFERENCE 3: 134:158493

REFERENCE 4: 133:295360

REFERENCE 5: 128:292989

L5 ANSWER 25 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN 206205-21-2 REGISTRY

CN Antigen (Babesia microtia 267-amino acid) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 247: PN: WO0185947 SEQID: 28 claimed protein

CN 28: PN: US6183976 SEQID: 28 claimed protein

CN 28: PN: US6214971 SEQID: 28 claimed protein

CN 28: PN: WO0060090 SEQID: 28 unclaimed protein

CN Antigen BMNI-13 (Babesia microti)

CN Antigen BMNI-13 (Babesia microtia fragment)

CN Antigen BMNI-13 (Bombesia microtia antigen BMNI-17 fragment)

CI MAN

SQL 267

```
SEQ      1 LWFIKMVSFK SILVPYITLF LMSGAVFASD TDPEAGGPSG TVGPSEAGGP
          =====
          51 SEAGGPSGTG WPSEAGGPSE AGGPSGTGWP SEAGWSSERF GYQLLPYSRR
          =====
          101 IVTFNEVCLS YIYKHSVMIL ERDRVNDGHK DYIEEKTKEK NKLKKELEKC
          151 FPEQYSLMKK EELARIFDNA STISSKYKLL VDEISNKAYG TLEGPAADNF
          201 DHFRNIWKS I VLKDMFIYCD LLLQHLYYKF YYDNTINDIK KNFDESKSKA
          251 LVL RDKITKK DVYVNDH
```

HITS AT: 34-87

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

REFERENCE 3: 134:158493

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

REFERENCE 6: 128:292989

L5 ANSWER 26 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN 206205-20-1 REGISTRY

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CN Antigen (Babesia microtia 121-amino acid) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 246: PN: WO0185947 SEQID: 27 claimed protein
CN 27: PN: US6183976 SEQID: 27 claimed protein
CN 27: PN: US6214971 SEQID: 27 claimed protein
CN 27: PN: WO0060090 SEQID: 27 unclaimed protein
CN Antigen BMNI-12 (Babesia microti)
CN Antigen BMNI-12 (Babesia microtia fragment)
CN Antigen BMNI-12 (Bombesia microtia antigen BMNI-17 fragment)
CI MAN
SQL 121

SEQ 1 LWFIKMVSFK SILVPYITLF LMSGAVFASD TDPEAGGPSE AGGPSEAGGP
51 SGTVGPSEAG GPSEAGGPSG TGWPSEAGGP SEAGGPSGTG WPSEAGWSSE
101 RFGYQLLPYS RRIVTFNEVC L
HITS AT: 34-99

REFERENCE 1: 135:367756
REFERENCE 2: 134:294507
REFERENCE 3: 134:158493
REFERENCE 4: 133:295360
REFERENCE 5: 131:85409
REFERENCE 6: 128:292989

L5 ANSWER 27 OF 32 REGISTRY COPYRIGHT 2002 ACS
RN 206205-16-5 REGISTRY
CN Antigen (Babesia microtia 303-amino acid) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 23: PN: US6183976 SEQID: 23 claimed protein
CN 23: PN: US6214971 SEQID: 23 claimed protein
CN 23: PN: WO0060090 SEQID: 23 unclaimed protein
CN 242: PN: WO0185947 SEQID: 23 claimed protein
CN Antigen BMNI-6 (Babesia microti)
CN Antigen BMNI-6 (Babesia microtia fragment)
CN Antigen BMNI-6 (Bombesia microtia antigen BMNI-17 fragment)
CI MAN
SQL 303

SEQ 1 LWFIKMVSFK SILVPYITLF LMSGAVFAGD TDREAGGPSG TVGPSEAGGP
51 SEAGGPSEAG GPSEAGGPSE AGGPSEAGGP SEAGGPSEAG GPSGTGWPSE
101 AGWPSEAGWP SEAGWPSEAG WPSEAGWPSE RFGYQLLWYS RRIVIFNEIY
151 LSHIYHSVM ILERDRVNDG HKDYIEEKT EKNKLKKELE KCFPEQYSLM
201 KKEELARIID NASTISSKYK LLVDEISNKA YGTLEGPAAD DFDHFRNIWK
251 SIVPKNMFY CDLLKHLIR KFYCDNTIND IKKNFDDIEK LGCFQARSFL
301 PVN
HITS AT: 34-129

REFERENCE 1: 135:367756

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09/853079

REFERENCE 2: 134:294507

REFERENCE 3: 134:158493

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

REFERENCE 6: 128:292989

L5 ANSWER 28 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN 206205-13-2 REGISTRY

CN Antigen (Babesia microtia 367-amino acid) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 20: PN: US6183976 SEQID: 20 claimed protein

CN 20: PN: US6214971 SEQID: 20 claimed protein

CN 20: PN: WO0060090 SEQID: 20 unclaimed protein

CN 239: PN: WO0185947 SEQID: 20 claimed protein

CN Antigen BMNI-3 (Babesia microti)

CN Antigen BMNI-3 (Babesia microtia fragment)

CN Antigen BMNI-3 (Babesia microtia antigen BMNI-17 fragment)

CI MAN

SQL 367

SEQ 1 LWFIKMVSFK SILVPYITLF LMSGAVFASD TDPEAGGPSE AGGPGSGTVGP

51 SEAGGPSEAG GPSGTGWPSG AGGPSEAGGP SEAGGPSEAG GPSGTGWPSG

101 TGWPSEAGWS SERFGYQLLP YSRIVIFNE VCLSYIYKHS VMILERDRVN

151 DGHKDYIEEK TKEKNKLKKE LEKCFPEQYS LMKKEELARI FDNASTISSK

201 YKLLVDEISN KAYGTLEGPA ADNFDHFRNI WKSIVLKD MF IYCDLLLQHL

251 IYKFYYDNTV NDIKKNFDES KSKALVLRDK ITKKG DYNT HFEDMIKELN

301 SAAEEFNKIV DIMISNIGDY DEYDSIASFK PFLSMITEIT KITKVS NVII

351 PGIKALTITV FLIFITK

HITS AT: 34-111

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

REFERENCE 3: 134:158493

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

REFERENCE 6: 128:292989

L5 ANSWER 29 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN 206205-12-1 REGISTRY

CN Antigen (Babesia microtia 310-amino acid) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 19: PN: US6183976 SEQID: 19 claimed protein

CN 19: PN: US6214971 SEQID: 19 claimed protein

CN 19: PN: WO0060090 SEQID: 19 unclaimed protein

CN 238: PN: WO0185947 SEQID: 19 claimed protein

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CN Antigen BMNI-2 (Babesia microti)
CN Antigen BMNI-2 (Babesia microtia fragment)
CN Antigen BMNI-2 (Bombesia microtia antigen BMNI-17 fragment)
CI MAN
SQL 310

SEQ 1 MSGAVFASDT DPEAGGPSEA GGPSGTVGPS EAGGPSEAGG PSGTVGPSEA
=====
51 GGPSEAGGPS GTGWPSEAGG PSEAGGPSGT VGPSEAGGPS EAGGPSGTGW
=====
101 PSEAGGPSEA GGPSEAGGPS EAGGPSGTGW PSGTGWPSSEA GWSSERFGYQ
=====
151 LLPYSRRIVI FNEVCLSYIY KHSVMIERD RVNDGHKDYI EEKTKEKNKL
201 KKELEKCFPE QYSLMKKEEL ARIFDNASTI SSKYKLLVDE ISNKAYGTLE
251 GPAADNFDHF RNIWKSIVLK DMFIYCDLLL QHLYKFYYD NTVNDIKKNF
301 DESWTQTLKE

HITS AT: 13-144

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

REFERENCE 3: 134:158493

REFERENCE 4: 133:295360

REFERENCE 5: 131:85409

REFERENCE 6: 128:292989

L5 ANSWER 30 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN 206205-11-0 REGISTRY

CN Antigen (Babesia microtia 263-amino acid) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 18: PN: US6183976 SEQID: 18 claimed protein
CN 18: PN: US6214971 SEQID: 18 claimed protein
CN 18: PN: WO0060090 SEQID: 18 unclaimed protein
CN 237: PN: WO0185947 SEQID: 18 claimed protein

CN Antigen BMNI-1 (Babesia microti)

CN Antigen BMNI-1 (Babesia microtia fragment)

CN Antigen BMNI-1 (Bombesia microtia antigen BMNI-17 fragment)

CI MAN

SQL 263

SEQ 1 LFLMSGAVFA SDTDPEAGGP SEAGGPSGTV GPSEAGGPSE AGGPSGTGWP
=====
51 SEAGGPSEAG GPSEAGGPSE AGGPSGTGWP SGTGWPSEAG WSSERFGYQL
=====
101 LPYSRRIVIF NEVCLSYIYK HSVMIERDR VNDGHKDYIE EKTKEKNKLK
151 KELEKCFPEQ YSLMKKEELA RIFDNASTIS SKYKLLVDEI SNKAYGTLEG
201 PAADNFDHFR NIWKSIVLKD MFIYCDLLLQ HLIYKFYYDN TVNDIKKNFD
251 ESKSKALVLR DKI

HITS AT: 16-93

REFERENCE 1: 135:367756

REFERENCE 2: 134:294507

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09/853079

REFERENCE 3: 134:158493
REFERENCE 4: 133:295360
REFERENCE 5: 131:85409
REFERENCE 6: 128:292989

L5 ANSWER 31 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN 205488-54-6 REGISTRY

CN L-Serine, L-.alpha.-glutamyl-L-alanylglycylglycyl-L-prolyl-L-serylglycyl-L-threonyl-L-valylglycyl-L-prolyl-L-serylglycyl-L-threonylglycyl-L-tryptophyl-L-prolyl-L-seryl-L-.alpha.-glutamyl-L-alanylglycyl-L-tryptophylglycyl-L-seryl-L-.alpha.-glutamyl-L-alanylglycyl-L-tryptophyl-L-seryl- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 17: PN: WO0185947 SEQID: 48 unclaimed sequence
CN 48: PN: US6183976 SEQID: 48 claimed sequence
CN 48: PN: US6214971 SEQID: 48 unclaimed sequence
CN 89: PN: WO0060090 SEQID: 48 unclaimed sequence
CN Antigen BABS-2 (Babesia microtia fragment)
SQL 30

SEQ 1 EAGGPSGTVG PSGTGWPSEA GWGSEAGWSS

=====

HITS AT: 1-18, 25-30

REFERENCE 1: 135:367756
REFERENCE 2: 134:294507
REFERENCE 3: 134:158493
REFERENCE 4: 133:295360
REFERENCE 5: 131:85409
REFERENCE 6: 128:292989

L5 ANSWER 32 OF 32 REGISTRY COPYRIGHT 2002 ACS

RN 205488-48-8 REGISTRY

CN L-Serine, L-seryl-L-.alpha.-glutamyl-L-alanylglycylglycyl-L-prolyl-L-seryl-L-.alpha.-glutamyl-L-alanylglycylglycyl-L-prolyl-L-serylglycyl-L-threonylglycyl-L-tryptophyl-L-threonyl-L-serylglycyl-L-threonylglycyl-L-tryptophyl-L-prolyl-L-seryl-L-.alpha.-glutamyl-L-alanylglycyl-L-tryptophyl- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 16: PN: WO0185947 SEQID: 47 unclaimed sequence
CN 47: PN: US6183976 SEQID: 47 claimed sequence
CN 47: PN: US6214971 SEQID: 47 unclaimed sequence
CN 88: PN: WO0060090 SEQID: 47 unclaimed sequence
CN Antigen BABS-1 (Babesia microtia fragment)
SQL 30

SEQ 1 SEAGGPSEAG GPSGTGWTSG TGWPSEAGWS

=====

HITS AT: 2-13, 20-25

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09/853079

REFERENCE 1: 135:367756
REFERENCE 2: 134:294507
REFERENCE 3: 134:158493
REFERENCE 4: 133:295360
REFERENCE 5: 131:85409
REFERENCE 6: 128:292989

(FILE 'HCAPLUS, MEDLINE, BIOSIS, EMBASE, WPIDS, CONFSCI, SCISEARCH,
JICST-EPLUS, JAPIO' ENTERED AT 12:09:33 ON 30 AUG 2002)

L6 4373 S "REED S"?/AU

- Author(s)

L8 215 SEA ABB=ON PLU=ON "LODES M"?/AU
L9 927 SEA ABB=ON PLU=ON "HOUGHTON R"?/AU
L10 177 SEA ABB=ON PLU=ON "SLEATH P"?/AU
L11 295 SEA ABB=ON PLU=ON "MCNEILL P"?/AU
L12 416 SEA ABB=ON PLU=ON "HOMER M"?/AU
L13 113 SEA ABB=ON PLU=ON "SECRIST H"?/AU
L14 2 SEA ABB=ON PLU=ON L6 AND L8 AND L9 AND L10 AND L11 AND
L12 AND L13
L15 231 SEA ABB=ON PLU=ON L6 AND (L8 OR L9 OR L10 OR L11 OR
L12 OR L13)
L16 115 SEA ABB=ON PLU=ON L8 AND (L9 OR L10 OR L11 OR L12 OR
L13)
L17 85 SEA ABB=ON PLU=ON L9 AND (L10 OR L11 OR L12 OR L13)
L18 19 SEA ABB=ON PLU=ON L10 AND (L11 OR L12 OR L13)
L19 2 SEA ABB=ON PLU=ON L11 AND (L12 OR L13)
L20 2 SEA ABB=ON PLU=ON L12 AND L13
L21 43 SEA ABB=ON PLU=ON (L15 OR L16 OR L17 OR L6 OR L8 OR L9
OR L10 OR L11 OR L12 OR L13) AND MICROTI

~~L22~~ 56 SEA ABB=ON PLU=ON L14 OR L18 OR L19 OR L20 OR L21

~~L23~~ 27 DUP REM L22 (29 DUPLICATES REMOVED)

L23 ANSWER 1 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 1
ACCESSION NUMBER: 2002:275811 HCAPLUS
DOCUMENT NUMBER: 136:308523
TITLE: Compositions and methods for WT1 specific
immunotherapy
INVENTOR(S): Gaiger, Alexander; McNeill, Patricia D.
; Smithgall, Molly; Moulton, Gus; Vedvick,
Thomas S.; Sleath, Paul R.; Mossman,
Sally; Evans, Lawrence; Spies, A. Gregory;
Boydston, Jeremy
PATENT ASSIGNEE(S): Corixa Corporation, USA
SOURCE: PCT Int. Appl., 260 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002028414	A1	20020411	WO 2001-US31139	20011003

Searcher : Shears 308-4994

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09/853079

WO 2002028414 B1 20020718

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,
GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG,
KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH,
CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE,
TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,
TD, TG

PRIORITY APPLN. INFO.:

US 2000-684361 A 20001006
US 2000-685830 A 20001009
US 2001-785019 A 20010215
US 2001-938864 A 20010824

AB Compns. and methods for the therapy of malignant diseases, such as leukemia and cancer, are disclosed. The compns. comprise one or more of a WT1 polynucleotide, a WT1 polypeptide, an antigen-presenting cell presenting a WT1 polypeptide, an antibody that specifically binds to a WT1 polypeptide; or a T cell that specifically reacts with a WT1 polypeptide. Such compns. may be used, for example, for the prevention and treatment of metastatic diseases.

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 2 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 2

ACCESSION NUMBER: 2001:833517 HCAPLUS

DOCUMENT NUMBER: 135:367756

TITLE: Babesia **microti** antigens and methods for the diagnosis and treatment of Babesia **microti** infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton, Raymond L.; Sleath, Paul R.; McNeill, Patricia D.; Homer, Mary J.; Secrist, Heather

PATENT ASSIGNEE(S): Corixa Corporation, USA

SOURCE: PCT Int. Appl., 195 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 7

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001085947	A2	20011115	WO 2001-US15192	20010509
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH,			

Searcher : Shears 308-4994

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09/853079

CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE,
TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD,
TG

US 2001029295 A1 20011011 US 2000-737178 20001213
PRIORITY APPLN. INFO.: US 2000-569098 A 20000510
US 2000-605724 A 20000627
US 2000-656688 A 20000907
US 2000-685436 A 20001010
US 2000-737178 A 20001213
US 2001-794764 A 20010226
US 1996-723142 A2 19961001
US 1997-845258 A2 19970424
US 1997-990571 A2 19971211
WO 1998-US26437 A2 19981211
US 1999-286488 A2 19990405
US 2000-528784 A2 20000317
WO 2000-US9136 A2 20000405

AB Compds. and methods for the diagnosis and treatment of B.
microti infection are disclosed. The compds. provided
include polypeptides that contain at least one antigenic portion of
a B. **microti** antigen and DNA sequences encoding such
polypeptides. Antigenic epitopes of such antigens are also
provided, together with pharmaceutical compns. and immunogenic
compns. comprising such polypeptides, DNA sequences or antigenic
epitopes. Diagnostic kits contg. such polypeptides, DNA sequences
or antigenic epitopes and a suitable detection reagent may be used
for the detection of B. **microti** infection in patients and
biol. samples. Antibodies directed against such polypeptides and
antigenic epitopes are also provided.

L23 ANSWER 3 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 3
ACCESSION NUMBER: 2001:748300 HCAPLUS
DOCUMENT NUMBER: 135:299589
TITLE: Nucleic acids and proteins for the diagnosis and
treatment of Babesia **microti** infection
INVENTOR(S): Reed, Steven G.; Lodes, Michael
J.; Houghton, Raymond L.;
Sleath, Paul R.; McNeill, Patricia
D.
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 26 pp., Cont.-in-part of
U. S. Ser. No. 685,436.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 7
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2001029295	A1	20011011	US 2000-737178	20001213
US 6306396	B1	20011023	US 1996-723142	19961001
US 6183976	B1	20010206	US 1997-845258	19970424
US 6214971	B1	20010410	US 1997-990571	19971211
WO 9929869	A1	19990617	WO 1998-US26437	19981211
W: AU, CA, JP, MX, NZ				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				

Searcher : Shears 308-4994

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09/853079

WO 2000060090	A1	20001012	WO 2000-US9136	20000405
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN,			
	CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,			
	HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,			
	LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,			
	RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,			
	US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,			
	DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF,			
	BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			

WO 2001085947	A2	200111115	WO 2001-US15192	20010509
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.:

US 1996-723142	A2 19961001
US 1997-845258	A2 19970424
US 1997-990571	A2 19971211
WO 1998-US26437	A2 19981211
US 1999-286488	A2 19990405
US 2000-528784	A2 20000317
WO 2000-US9136	A2 20000405
US 2000-569098	A2 20000510
US 2000-605724	A2 20000627
US 2000-656688	A2 20000907
US 2000-685436	A2 20001010
US 2000-737178	A 20001213
US 2001-794764	A 20010226

AB Compsds. and methods for the diagnosis and treatment of B. microtiinfection are disclosed. The compds. provided include polypeptides that contain at least one antigenic portion of a B. microti antigen and DNA sequences encoding such polypeptides. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compns. and immunogenic compns. comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of B. microtiinfection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

L23 ANSWER 4 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 4
ACCESSION NUMBER: 2001:255942 HCAPLUS
DOCUMENT NUMBER: 134:294507
TITLE: Compounds and methods for the diagnosis and
treatment of Babesia **microti** infection
INVENTOR(S): **Reed, Steven G.; Lodes, Michael**
J.; Houghton, Raymond
PATENT ASSIGNEE(S): Corixa Corporation, USA
SOURCE: U.S., 78 pp., Cont.-in-part of U.S. Ser. No.
845,258.
CODEN: USXXAM

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09/853079

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 7
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6214971	B1	20010410	US 1997-990571	19971211
US 6306396	B1	20011023	US 1996-723142	19961001
US 6183976	B1	20010206	US 1997-845258	19970424
WO 9929869	A1	19990617	WO 1998-US26437	19981211
W: AU, CA, JP, MX, NZ				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
AU 9918204	A1	19990628	AU 1999-18204	19981211
US 2001029295	A1	20011011	US 2000-737178	20001213

PRIORITY APPLN. INFO.:

US 1996-723142	A2	19961001
US 1997-845258	A2	19970424
US 1997-990571	A	19971211
WO 1998-US26437	W	19981211
US 1999-286488	A2	19990405
US 2000-528784	A2	20000317
WO 2000-US9136	A2	20000405
US 2000-569098	A2	20000510
US 2000-605724	A2	20000627
US 2000-656688	A2	20000907
US 2000-685436	A2	20001010

AB Compds. and methods for the diagnosis and treatment of Babesia **microti** infection are disclosed. The compds. provided include polypeptides that contain at least one antigenic portion of a B. **microti** antigen and DNA sequences encoding such polypeptides. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compns. and vaccines comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of B. **microti** infection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

REFERENCE COUNT: 13 THERE ARE 13 CITED REFERENCES AVAILABLE
FOR THIS RECORD. ALL CITATIONS AVAILABLE
IN THE RE FORMAT

L23 ANSWER 5 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 5
ACCESSION NUMBER: 2001:91448 HCAPLUS
DOCUMENT NUMBER: 134:158493
TITLE: Nucleic acids and proteins for the diagnosis and
treatment of Babesia **microti** infection
INVENTOR(S): Reed, Steven G.; Lodes, Michael
J.; Houghton, Raymond;
Sleath, Paul R.
PATENT ASSIGNEE(S): Corixa Corporation, USA
SOURCE: U.S., 66 pp., Cont.-in-part of U.S. Ser. No.
723,142.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 7

Searcher : Shears 308-4994

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09/853079

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6183976	B1	20010206	US 1997-845258	19970424
US 6306396	B1	20011023	US 1996-723142	19961001
EP 834567	A2	19980408	EP 1997-117067	19971001
EP 834567	A3	19990331		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
US 6214971	B1	20010410	US 1997-990571	19971211
US 2001029295	A1	20011011	US 2000-737178	20001213

PRIORITY APPLN. INFO.:

US 1996-723142	A2	19961001
US 1997-845258	A	19970424
US 1997-990571	A2	19971211
WO 1998-US26437	A2	19981211
US 1999-286488	A2	19990405
US 2000-528784	A2	20000317
WO 2000-US9136	A2	20000405
US 2000-569098	A2	20000510
US 2000-605724	A2	20000627
US 2000-656688	A2	20000907
US 2000-685436	A2	20001010

OTHER SOURCE(S): MARPAT 134:158493

AB Compds. and methods for the diagnosis and treatment of Babesia **microti** infection are disclosed. The compds. provided include polypeptides that contain at least one antigenic portion of a B. **microti** antigen and DNA sequences encoding such polypeptides. The DNA sequences encoding B. **microti** antigens were prep'd. by screening a B. **microti** expression library with sera obtained from patients infected with B. **microti**. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compns. and vaccines comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of B. **microti** infection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 6 OF 27 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 2002:6260 BIOSIS

DOCUMENT NUMBER: PREV200200006260

TITLE: Compounds and methods for the diagnosis and treatment of B. **microti** infection.

AUTHOR(S): Reed, Steven G.; Lodes, Michael J.
; Houghton, Raymond; Sleath, Paul
R.

ASSIGNEE: Corixa Corporation

PATENT INFORMATION: US 6306396 October 23, 2001

SOURCE: Official Gazette of the United States Patent and Trademark Office Patents, (Oct. 23, 2001) Vol. 1251, No. 4, pp. No Pagination. e-file.
ISSN: 0098-1133.

DOCUMENT TYPE: Patent

Searcher : Shears 308-4994

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09/853079

LANGUAGE: English

AB Compounds and methods for the diagnosis and treatment of B. **microti** infection are disclosed. The compounds provided include polypeptides that contain at least one antigenic portion of a B. **microti** antigen and DNA sequences encoding such polypeptides. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compositions and vaccines comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits containing such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of B. **microti** infection in patients and biological samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

L23 ANSWER 7 OF 27 WPIDS (C) 2002 THOMSON DERWENT

ACCESSION NUMBER: 2002-062250 [08] WPIDS

CROSS REFERENCE: 1998-609891 [51]; 2000-160675 [11]

DOC. NO. NON-CPI: N2002-046184

DOC. NO. CPI: C2002-017807

TITLE: Novel polynucleotide encoding polypeptides useful for detecting Ehrlichia infection in patients and biological samples, and for treating human granulocytic ehrlichiosis, comprise an Ehrlichia antigen.

DERWENT CLASS: B04 C06 D16 S03

INVENTOR(S): HOUGHTON, R L; LODES, M J;

MCNEILL, P D; REED, S G

PATENT ASSIGNEE(S): (CORI-N) CORIXA CORP; (HOUG-I) HOUGHTON R L;
(LODE-I) LODES M J; (MCNE-I) MCNEILL P D; (REED-I) REED S G

COUNTRY COUNT: 95

PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 2001085949	A2	20011115	(200208)*	EN	132
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW MZ NL OA PT SD SE SL SZ TR TZ UG ZW					
W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ					
DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE					
KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO					
NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ					
VN YU ZA ZW					
AU 2001059507	A	20011120	(200219)		
US 2002068343	A1	20020606	(200241)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 2001085949	A2	WO 2001-US14518	20010504
AU 2001059507	A	AU 2001-59507	20010504
US 2002068343	A1	CIP of US 1997-821324	19970321
		CIP of US 1997-975762	19971120
		CIP of US 1998-106582	19980629
		CIP of US 1998-159469	19980923
		CIP of US 1999-295028	19990420
		CIP of US 2000-566617	20000508

Searcher : Shears 308-4994

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CIP of US 2000-693542 20001020
US 2001-798042 20010302

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 2001059507	A	Based on WO 200185949
US 2002068343	A1	CIP of US 6207169
		CIP of US 6231869
		CIP of US 6277381
		CIP of US 6306402

PRIORITY APPLN. INFO: US 2001-798042 20010302; US 2000-566617
20000508; US 2000-693542 20001020; US
1997-821324 19970321; US 1997-975762
19971120; US 1998-106582 19980629; US
1998-159469 19980923; US 1999-295028 19990420

AN 2002-062250 [08] WPIDS
CR 1998-609891 [51]; 2000-160675 [11]
AB WO 200185949 A UPAB: 20020701

NOVELTY - An isolated polynucleotide (I) comprising a sequence (S1) chosen from 36 nucleotides of defined bp fully in the specification such as 1345, 1132, 554, 559, 201, 467, 530, 1185, 1131, 800, 1011, 513, 464, 527, 464, 860, 484 or 1039 bp, complement of (S1), a sequence hybridizable under moderate stringent conditions to (S1), a sequence which is 75% or 90% identical to (S1) or degenerate variant of (S1), is new.

DETAILED DESCRIPTION - (I) is chosen from the determined DNA sequence of human granulocytic ehrlichiosis (HGE)-1, HGE-3, HGE-6, HGE-8, HGE-11-13, HGE-23, HGE-24, the 5' DNA sequence of HGE-7, HGE-2, HGE-9, HGE-14-18, HGE-25, extended DNA sequences of HGE-2, HGE-7, HGE-8, HGE-11, HGE-14-16, HGE-18, HGE-23, HGE-25, the determined 3' DNA sequence of HGE-17, the full-length cDNA sequence for HGE-17, corrected cDNA sequence for HGE-14, HGE-1, and the reverse complement of HGE-2, HGE-14 and HGE-15. INDEPENDENT CLAIMS are also included for the following:

- (1) an isolated polypeptide (II) comprising a sequence encoded by (I) or a sequence 70% or 90% identical to a sequence encoded by (I);
- (2) an isolated antigenic epitope (III) of an Ehrlichia antigen comprising a sequence of 41 or 125 amino acids defined in the specification;
- (3) an isolated polypeptide comprising at least two antigenic epitopes as above;
- (4) a recombinant expression vector (IV) comprising (I);
- (5) a host cell (V) transformed with (IV);
- (6) a fusion protein (VI) comprising (II) or (III);
- (7) a diagnostic kit comprising (II), (III) and (VI) and a detection reagent;
- (8) a diagnostic kit comprising at least two oligonucleotide probes or primer specific for (I);
- (9) an isolated antibody (VII), or its antigen binding fragment that specifically binds to (II) or (III);
- (10) a composition (VIII) comprising any one chosen from (I)-(III), (VI) and (VII), and immunostimulants;
- (11) detecting Ehrlichia infection in a biological sample, by contacting the sample with oligonucleotide primers or probes, or a

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binding agent capable of binding to (II), (III) or (VI) and detecting in the sample the presence of amplified polynucleotide sequence, polynucleotide sequence that hybridizes to the probe or polypeptide that binds to the binding agent; and

(12) detecting a disorder of Ehrlichia infection, Lyme disease and *B. microti* infection in a patient, by contacting the biological sample with (II), a Lyme disease antigen, and a *B. microti* antigen, and detecting the presence of antibodies in the biological sample that bind to either the polypeptide, Lyme disease antigen or the *B. microti* antigen.

ACTIVITY - Antibiotic. No supporting data is provided.

MECHANISM OF ACTION - Vaccine. No supporting data is given.

USE - (II), (III) and (VI) are useful for detecting Ehrlichia infection in a patient. (VIII) is useful for stimulating an immune response in a patient, for treating Ehrlichia infection in a patient (claimed). (II) is useful for serodiagnosis and treatment of human granulocytic ehrlichiosis (HGE). (VII) is useful in diagnostic test to detect the presence of Ehrlichia antigens and for detecting Ehrlichia infection in a patient.

Dwg.0/2

L23 ANSWER 8 OF 27 SCISEARCH COPYRIGHT 2002 ISI (R)
 ACCESSION NUMBER: 2001:943909 SCISEARCH
 THE GENUINE ARTICLE: 494QP
 TITLE: Innate resistance to Babesia infection is influenced by genetic background and gender
 AUTHOR: Aguilar-Delfin I; **Homer M J**; Wettstein P J; Persing D H (Reprint)
 CORPORATE SOURCE: Corixa Corp, Suite 200, 1124 Columbia St, Seattle, WA 98104 USA (Reprint); Mayo Clin & Mayo Fdn, Dept Immunol, Rochester, MN 55905 USA; Corixa Corp, Seattle, WA 98104 USA; Infect Dis Res Inst, Seattle, WA 98104 USA
 COUNTRY OF AUTHOR: USA
 SOURCE: INFECTION AND IMMUNITY, (DEC 2001) Vol. 69, No. 12, pp. 7955-7958.
 Publisher: AMER SOC MICROBIOLOGY, 1752 N ST NW, WASHINGTON, DC 20036-2904 USA.
 ISSN: 0019-9567.
 DOCUMENT TYPE: Article; Journal
 LANGUAGE: English
 REFERENCE COUNT: 30

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

AB Infection of severe combined immunodeficient mice with Babesia sp. strain WA1 was studied to assess the contributions of innate and adaptive immunity in resistance to acute babesiosis. The scid mutation showed little effect in genetically susceptible C3H mice and did not decrease the inherent resistance of C57BL/6 mice to the infection, suggesting that innate immunity plays a central role in determining the course of Babesia infection in these strains. In contrast, the scid mutation dramatically impaired resistance in moderately susceptible BALB/c mice, suggesting that acquired immunity may play an important secondary role. In comparison to their female counterparts, male mice of different genetic backgrounds showed increased resistance to the infection, indicating that the gender of the host may influence protection against babesiosis.

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L23 ANSWER 9 OF 27 SCISEARCH COPYRIGHT 2002 ISI (R)
ACCESSION NUMBER: 2001:546315 SCISEARCH
THE GENUINE ARTICLE: 447RN
TITLE: Serodiagnosis of human granulocytic ehrlichiosis by
using novel combinations of immunoreactive
recombinant proteins
AUTHOR: **Lodes M J (Reprint)**; Mohamath R; Reynolds
L D; **McNeill P**; Kolbert C P; Bruinsma E S;
Benson D R; Hofmeister E; **Reed S G**;
Houghton R L; Persing D H
CORPORATE SOURCE: Corixa Corp, 1124 Columbia St, Suite 200, Seattle,
WA 98104 USA (Reprint); Infect Dis Res Inst,
Seattle, WA 98104 USA; Corixa Corp, Seattle, WA
98104 USA; Univ Washington, Dept Pathobiol, Seattle,
WA 98195 USA; Mayo Clin & Mayo Fdn, Dept Lab Med &
Pathol, Rochester, MN 55905 USA
COUNTRY OF AUTHOR: USA
SOURCE: JOURNAL OF CLINICAL MICROBIOLOGY, (JUL 2001) Vol.
39, No. 7, pp. 2466-2476.
Publisher: AMER SOC MICROBIOLOGY, 1752 N ST NW,
WASHINGTON, DC 20036-2904 USA.
ISSN: 0095-1137.
DOCUMENT TYPE: Article; Journal
LANGUAGE: English
REFERENCE COUNT: 44

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

AB A panel of seven recombinant antigens, derived from Ehrlichia phagocytophila (the agent of human granulocytic ehrlichiosis), was evaluated by class-specific enzyme-linked immunosorbent assays (ELISAs) for utility in the diagnosis of the infection. Fourteen genomic fragments, obtained by serologic expression screening, contained open reading frames (ORFs) encoding 16 immunodominant antigens. Eleven of these antigens were members of the major surface protein (MSP) multigene family. Alignment of their predicted protein sequences revealed a pattern of conserved sequences, which contained short direct repeats, flanking a variable region. In addition, two genomic clones contained two and three MSP ORFs, respectively, indicating that these genes are clustered in tandem copies. The implications for this pattern of both genomic and protein arrangements in antigenic variations of MSPs and in their utilities in a diagnostic assay are discussed. In addition to two MSP recombinant antigens (rHGE-1 and -3) and a fusion protein of these antigens (rErf-1), five further recombinants were evaluated by ELISA. Two of these antigens (rHGE-14 and -15) were novel, while a third (rHGE-2), with no known function, has been described. The final two recombinant antigens (rHGE-9 and -17) represent overlapping segments of the ankyrin gene (ank). The addition of rHGE-9 ELISA data resulted in the detection of 78% (21 of 27) of acute-phase sera. When serologic data for all recombinants are combined, 96.2% (26 of 27) of convalescent-phase patient serum samples and 85.2% (23 of 27) of acute-phase patient serum samples are detected, indicating the potential of these antigens for use in the development of a rapid serologic assay for the detection of E. phagocytophila infection.

L23 ANSWER 10 OF 27 SCISEARCH COPYRIGHT 2002 ISI (R)
ACCESSION NUMBER: 2001:818201 SCISEARCH
THE GENUINE ARTICLE: 472TV

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09/853079

TITLE: Peptide ELISA for detection of antibodies to Babesia
microti in serum.
AUTHOR: **Houghton R L (Reprint); Homer M J**
; Reynolds L D; **Sleath P C**; Cable R G;
Militscher J E; **Lodes M J**; Berardi V;
Leiby D A; Persing D H
CORPORATE SOURCE: Corixa Corp, Seattle, WA USA; Amer Red Cross,
Farmington, CT USA; Amer Red Cross, Rockville, MD
USA; Imugen, Norwood, MA USA
COUNTRY OF AUTHOR: USA
SOURCE: TRANSFUSION, (SEP 2001) Vol. 41, No. 9, Supp. [S],
pp. 13S-13S.
Publisher: AMER ASSOC BLOOD BANKS, 8101 GLENBROOK
RD, BETHESDA, MD 20814-2749 USA.
ISSN: 0041-1132.
DOCUMENT TYPE: Conference; Journal
LANGUAGE: English
REFERENCE COUNT: 0

L23 ANSWER 11 OF 27 SCISEARCH COPYRIGHT 2002 ISI (R)
ACCESSION NUMBER: 2001:818200 SCISEARCH
THE GENUINE ARTICLE: 472TV
TITLE: Evidence for transmission of Babesia **microti**
from Connecticut blood donors to recipients
AUTHOR: Cable R G (Reprint); Badon S; Trouem-Trend J;
Militscher J E; **Houghton R L; Lodes M**
J; Persing D H; Eberhard M L; Pleniazek N J;
Herwaldt B L; Leiby D A
CORPORATE SOURCE: Amer Red Cross Blood Serv, Farmington, CT USA; Amer
Red Cross, Holland Lab, Rockville, MD USA; Corixa
Corp, Seattle, WA USA; Ctr Dis Control & Prevent,
Atlanta, GA USA
COUNTRY OF AUTHOR: USA
SOURCE: TRANSFUSION, (SEP 2001) Vol. 41, No. 9, Supp. [S],
pp. 12S-13S.
Publisher: AMER ASSOC BLOOD BANKS, 8101 GLENBROOK
RD, BETHESDA, MD 20814-2749 USA.
ISSN: 0041-1132.
DOCUMENT TYPE: Conference; Journal
LANGUAGE: English
REFERENCE COUNT: 0

L23 ANSWER 12 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 6
ACCESSION NUMBER: 2000:725776 HCAPLUS
DOCUMENT NUMBER: 133:295360
TITLE: Antigens of Babesia microtia for use in the
diagnosis, prophylaxis, and treatment of
babesiosis
INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton,
Raymond L.; **Sleath, Paul R.**;
Mcneill, Patricia D.
PATENT ASSIGNEE(S): Corixa Corp., USA
SOURCE: PCT Int. Appl., 118 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 7
PATENT INFORMATION:

Searcher : Shears 308-4994

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PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000060090	A1	20001012	WO 2000-US9136	20000405
W:				
				AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN,
				CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
				HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
				LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
				RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,
				US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW:				GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
				DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF,
				BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
EP 1169455	A1	20020109	EP 2000-921771	20000405
R:				AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
				PT, IE, SI, LT, LV, FI, RO
US 2001029295	A1	20011011	US 2000-737178	20001213
PRIORITY APPLN. INFO.:			US 1999-286488	A 19990405
			US 2000-528784	A 20000317
			US 1996-723142	A2 19961001
			US 1997-845258	A2 19970424
			US 1997-990571	A2 19971211
			WO 1998-US26437	A2 19981211
			WO 2000-US9136	W 20000405
			US 2000-569098	A2 20000510
			US 2000-605724	A2 20000627
			US 2000-656688	A2 20000907
			US 2000-685436	A2 20001010

OTHER SOURCE(S): MARPAT 133:295360

AB Comps. and methods for the diagnosis and treatment of B. microtia infection are disclosed. The comps. provided include polypeptides that contain at least one antigenic portion of a B. microtia antigen and DNA sequences encoding such polypeptides. Antigenic epitopes of such antigens are also provided, together with pharmaceutical comps. and vaccines comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of B. microtia infection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided. Cloning of antigen genes by antibody screening of expression libraries with antiserum. Seventeen antigens were identified and several of these showed common sequences. The clones also contained telomere repeat sequences indicating that they were located near the telomere. Use of the antigens in diagnostic detection of B. microtia is demonstrated.

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 13 OF 27 WPIDS (C) 2002 THOMSON DERWENT

ACCESSION NUMBER: 2000-160675 [14] WPIDS

CROSS REFERENCE: 1998-609891 [51]; 2002-062250 [01]

DOC. NO. NON-CPI: N2000-119888

DOC. NO. CPI: C2000-050162

TITLE: New compounds and methods for the diagnosis of Ehrlichia infection, particularly Human granulocytic ehrlichiosis.

Searcher : Shears 308-4994

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DERWENT CLASS: A96 B04 D16 S03
INVENTOR(S): HOUGHTON, R L; LODES, M J;
MCNEILL, P D; REED, S G
PATENT ASSIGNEE(S): (CORI-N) CORIXA CORP; (HOUG-I) HOUGHTON R L;
(LODE-I) LODES M J; (MCNE-I) MCNEILL P D; (REED-I)
REED S G
COUNTRY COUNT: 85
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 2000000615	A2	20000106	(200014)*	EN	108
RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC					
MW NL OA PT SD SE SL SZ UG ZW					
W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI					
GB GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS					
LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK					
SL TJ TM TR TT UA UG UZ VN YU ZA ZW					
AU 9948474	A	20000117	(200026)		
US 6277381	B1	20010821	(200150)		
US 6306402	B1	20011023	(200165)		
EP 1144639	A2	20011017	(200169)	EN	
R: AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE					
US 2002064535	A1	20020530	(200240)		

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 2000000615	A2	WO 1999-US14793	19990629
AU 9948474	A	AU 1999-48474	19990629
US 6277381	B1 CIP of	US 1997-821324	19970321
	CIP of	US 1997-975762	19971120
	CIP of	US 1998-106582	19980629
	CIP of	US 1998-159469	19980923
		US 1999-295028	19990420
US 6306402	B1 CIP of	US 1997-821324	19970321
	CIP of	US 1997-975762	19971120
		US 1998-106582	19980629
EP 1144639	A2	EP 1999-932087	19990629
		WO 1999-US14793	19990629
US 2002064535	A1 CIP of	US 1997-821324	19970321
	CIP of	US 1997-975762	19971120
	Cont of	US 1998-106582	19980629
		US 1998-159469	19980923

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9948474	A Based on	WO 200000615
EP 1144639	A2 Based on	WO 200000615
US 2002064535	A1 CIP of	US 6207169
	CIP of	US 6231869
	Cont of	US 6306402

PRIORITY APPLN. INFO: US 1999-295028 19990420; US 1998-106582
19980629; US 1998-159469 19980923; US

Searcher : Shears 308-4994

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AN 2000-160675 [14] WPIDS
 CR 1998-609891 [51]; 2002-062250 [01]
 AB WO 200000615 A UPAB: 20020701

NOVELTY - A polypeptide (P) comprising an immunogenic portion of an Ehrlichia antigen or its variant that is encoded by one of 18 DNA sequences of 201-7091 base pairs (bp) (I)-(XVIII) (all sequences fully defined in the specification), their complements and DNA sequences that hybridize to sequences (I)-(XVIII), are new.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the following:

(1) An antigenic epitope (E) of an Ehrlichia antigen comprising an amino acid sequence of (XIX) or (XX) consisting of 41 and 125 amino acids (aa) respectively;

(2) A polypeptide (P') comprising at least two of (E);

(3) A DNA molecule (N) comprising a nucleotide sequence encoding (P) or (P');

(4) A recombinant expression vector (V) comprising (N);

(5) A host cell (H) transformed with (N);

(6) A fusion protein (F) comprising either at least one of (P) or (P') and/or at least one of (E) or a 376 aa sequence (XXI) and/or a 325 aa sequence (XXII).

(7) detecting Ehrlichia infection, Lyme disease and Babesia microti infection in a patient comprising:

(a) contacting a biological sample with at least one of (E), (P), (P') or (F) and a Lyme disease antigen and a B.microti antigen; and

(b) detecting the presence of antibodies that bind to (E), (P), (P') or (F) or the Lyme disease antigen or the B.microti antigen in the sample;

(8) A method similar to (7), comprising:

(a) contacting a biological sample with a specific binding agent to at least one of (E), (P), (P') or (F) or a Lyme disease antigen and a B.microti antigen; and

(b) detecting a polypeptide that binds to the binding agent, thereby detecting Ehrlichia infection;

(9) A method similar to (7), comprising:

(a) contacting the sample with one or more probe oligonucleotides (or at least two primer oligonucleotides in a PCR reaction) where at least one is specific for (N); and

(b) detecting in the sample a DNA sequence that hybridizes to (or amplifies in the presence of) the oligonucleotide primers, thereby detecting Ehrlichia infection;

(10) A diagnostic kit (K) comprising:

(a) at least one of (P), (P'), (E) or (F); and

(b) a detection agent;

(11) A diagnostic kit (K') comprising at least two oligonucleotide primers or one oligonucleotide probe whereby at least one is specific for (N);

(12) A monoclonal antibody or polyclonal antibody that binds to (P), (P') or (E); and

(13) Vaccines comprising at least one of (P), (P'), (N) or (E) and a non-specific immune enhancer such as an adjuvant;

(14) A polypeptide comprising an immunogenic portion of an Ehrlichia antigen associated with human granulocytic ehrlichiosis or its variant.

All sequences are fully defined in the specification.

USE - (P), (P'), (N), (F) and/or (E) are useful for the

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detection and treatment of Ehrlichia infection. (P), (P'), (F) and/or (E) can also be used to detect Lyme disease and B. microti infection. In particular, (P') can be used for the serodiagnosis and treatment of human granulocytic ehrlichiosis (HGE). Compositions of (P) or (P'), (N) and (E) can be used in the manufacture of a medicament for inducing protective immunity in a patient. The new vaccines are also used for inducing protective immunity in a patient.

ADVANTAGE - None given.

Dwg.0/2

L23 ANSWER 14 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 7
ACCESSION NUMBER: 2000:282570 HCAPLUS
DOCUMENT NUMBER: 133:72606
TITLE: Serological expression cloning of novel immunoreactive antigens of Babesia microti
AUTHOR(S): Lodes, Michael J.; Houghton, Raymond L.; Bruinsma, Elizabeth S.; Mohamath, Raodoh; Reynolds, Lisa D.; Benson, Darin R.; Krause, Peter J.; Reed, Steven G.; Persing, David H.
CORPORATE SOURCE: Corixa Corporation, Seattle, WA, 98104, USA
SOURCE: Infection and Immunity (2000), 68(5), 2783-2790
CODEN: INFIBR; ISSN: 0019-9567
PUBLISHER: American Society for Microbiology
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Increased recognition of the prevalence of human babesiosis in the United States, together with rising concern about the potential for transmission of this infection by blood transfusion, has provided motivation to develop definitive serol. and mol. tests for the causative agent, Babesia microti. To develop more sensitive and specific assays for B. microti, the authors screened a genomic expression library with patient serum pools. This screening resulted in the identification of three classes of novel genes and an addnl. two novel, unrelated genes, which together encode a total of 17 unique B. microti antigens. The first class (BMN1-2 family) of genes encodes seven closely related antigens with a degenerate six-amino-acid repeat that shows limited homol. to Plasmodium sp. merozoite and sporozoite surface antigens. A second class (BMN1-8 family) of genes encodes six related antigens, and the third class (BMN1-17 family) of genes encodes two related antigens. The two remaining genes code for novel and unrelated sequences. Among the three classes of antigens and remaining novel sequences, five were chosen to code for the most immunodominant antigens (BMN1-2, -9, -15, and -17 and MN-10). Western blot anal. with the resulting recombinant proteins indicated that these antigens were targets of humoral immune responses during B. microti infection in humans.

REFERENCE COUNT: 36 THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 15 OF 27 MEDLINE DUPLICATE 8
ACCESSION NUMBER: 2000344709 MEDLINE
DOCUMENT NUMBER: 20344709 PubMed ID: 10885987
TITLE: Babesiosis.
AUTHOR: Homer M J; Aguilar-Delfin I; Telford S R

Searcher : Shears 308-4994

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09/853079

CORPORATE SOURCE: 3rd; Krause P J; Persing D H
Corixa Corporation and The Infectious Disease
Research Institute, Seattle, Washington 98104, USA.
SOURCE: CLINICAL MICROBIOLOGY REVIEWS, (2000 Jul) 13 (3)
451-69. Ref: 245
Journal code: 8807282. ISSN: 0893-8512.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
General Review; (REVIEW)
(REVIEW, TUTORIAL)
LANGUAGE: English
FILE SEGMENT: Priority Journals
ENTRY MONTH: 200008
ENTRY DATE: Entered STN: 20000811
Last Updated on STN: 20000811
Entered Medline: 20000803

AB Babesiosis is an emerging, tick-transmitted, zoonotic disease caused by hematotropic parasites of the genus *Babesia*. Babesial parasites (and those of the closely related genus *Theileria*) are some of the most ubiquitous and widespread blood parasites in the world, second only to the trypanosomes, and consequently have considerable worldwide economic, medical, and veterinary impact. The parasites are intraerythrocytic and are commonly called piroplasms due to the pear-shaped forms found within infected red blood cells. The piroplasms are transmitted by ixodid ticks and are capable of infecting a wide variety of vertebrate hosts which are competent in maintaining the transmission cycle. Studies involving animal hosts other than humans have contributed significantly to our understanding of the disease process, including possible pathogenic mechanisms of the parasite and immunological responses of the host. To date, there are several species of *Babesia* that can infect humans, *Babesia microti* being the most prevalent. Infections with *Babesia* species generally follow regional distributions; cases in the United States are caused primarily by *B. microti*, whereas cases in Europe are usually caused by *Babesia divergens*. The spectrum of disease manifestation is broad, ranging from a silent infection to a fulminant, malaria-like disease, resulting in severe hemolysis and occasionally in death. Recent advances have resulted in the development of several diagnostic tests which have increased the level of sensitivity in detection, thereby facilitating diagnosis, expediting appropriate patient management, and resulting in a more accurate epidemiological description.

L23 ANSWER 16 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 9
ACCESSION NUMBER: 2000:74595 HCAPLUS
DOCUMENT NUMBER: 133:100272
TITLE: A polymorphic multigene family encoding an
immunodominant protein from *Babesia microti*
AUTHOR(S): Homer, M. J.; Bruinsma, E. S.;
Lodes, M. J.; Moro, M. H.; Telford, S.,
III; Krause, P. J.; Reynolds, L. D.; Mohamath,
R.; Benson, D. R.; Houghton, R. L.;
Reed, S. G.; Persing, D. H.
CORPORATE SOURCE: Department of Laboratory Medicine and Pathology,
Mayo Clinic, Rochester, MN, 55905, USA
SOURCE: Journal of Clinical Microbiology (2000), 38(1),

Searcher : Shears 308-4994

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09/853079

362-368

CODEN: JCMIDW; ISSN: 0095-1137

PUBLISHER: American Society for Microbiology

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Human babesiosis in the United States is caused predominantly by *Babesia microti*, a tick-transmitted blood parasite. Improved testing methods for the detection of infection with this parasite are needed, since asymptomatic *B. microti* infection represents a potential threat to the blood supply in areas where *B. microti* is endemic. We performed immunoscreening of an expression library of genomic DNA from a human isolate of *B. microti* (strain MN1). Among 17 unique immunoreactive clones, we identified 9 which represent a related family of genes with little sequence homol. to other known sequences but with an architecture resembling that of several surface proteins of *Plasmodium*. Within this family, a tandem array of a degenerate six-amino-acid repeat (SEAGGP, SEAGWP, SGTGWP, SGTVGP) was found in various lengths between relatively well conserved segments at the N and C termini. In order to examine within-clone variation, we developed a PCR protocol for direct recovery of a specific *bmnl-6* homolog directly from 30 human blood isolates, 4 corresponding hamster isolates, and 5 geog. corresponding *Peromyscus leucopus* (white-footed mouse) isolates. Isolates from the hamsters had the same sequences as those found in the corresponding human blood, suggesting that genetic variation of *bmnl-6* does not occur during passage. However, clones from different patients were often substantially different from each other with regard to the no. and location of the degenerate repeats within the *bmnl-6* homolog. Moreover, we found that strains that were closely related geog. were also closely related at the sequence level; nine patients, all from Nantucket Island, Mass., harbored clones that were indistinguishable from each other but that were distinct from those found in other northeastern or upper midwestern strains. We conclude that considerable genetic and antigenic diversity exists among isolates of *B. microti* from the United States and that geog. clustering of subtypes may exist. The nature of the *bmnl-6* gene family suggests a mechanism of antigenic variation in *B. microti* that may occur by recombination, differential expression, or a combination of both mechanisms.

REFERENCE COUNT: 43 THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 17 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 10

ACCESSION NUMBER: 2000:87236 HCAPLUS

DOCUMENT NUMBER: 133:57223

TITLE: Multiepitope synthetic peptide and recombinant protein for the detection of antibodies to *Trypanosoma cruzi* in patients with treated or untreated Chagas' disease

AUTHOR(S): Houghton, Raymond L.; Benson, Darin R.; Reynolds, Lisa; McNeill, Patricia; Sleath, Paul; Lodes, Michael; Skeiky, Yasir A. W.; Badaro, Roberto; Krettli, Antoniana U.; Reed, Steven G.

CORPORATE SOURCE: Corixa Corporation, Seattle, WA, 98104, USA
SOURCE: Journal of Infectious Diseases (2000), 181(1),

Searcher : Shears 308-4994

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325-330

CODEN: JIDIAQ; ISSN: 0022-1899

PUBLISHER: University of Chicago Press
DOCUMENT TYPE: Journal
LANGUAGE: English

AB A tetrapeptide and a recombinant protein, each representing 4 immunodominant epitopes of *Trypanosoma cruzi*, were tested by use of ELISA for the detection of serum antibodies. Sera from individuals with Chagas' disease, including persons untreated and successfully or unsuccessfully treated, were tested. These assays detected antibody in 100% of the parasitemias. The antibody reactivity decreased based on the success of treatment. Higher sensitivity was obsd. for tetrapeptide/recombinant protein assays than for lysate-based ELISA, and specificity was improved, particularly with *Leishmania* sera. The results indicate that multiepitope antigens provide a more sensitive and specific alternative to lysate for detection of anti-*T. cruzi* antibodies, as required for developing blood screening assays.

REFERENCE COUNT: 29 THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 18 OF 27 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 2000:394239 BIOSIS

DOCUMENT NUMBER: PREV200000394239

TITLE: Characterization of the *Babesia microti* chronic carrier state in a murine model.

AUTHOR(S): Homer, M. J. (1); Bruinsma, E. S. (1); Aguilar-Delfin, I. (1); Moro, M. J. (1); Persing, D. H.

CORPORATE SOURCE: (1) Mayo Fndn, Rochester, MN USA

SOURCE: Abstracts of the General Meeting of the American Society for Microbiology, (2000) Vol. 100, pp. 283. print.
Meeting Info.: 100th General Meeting of the American Society for Microbiology Los Angeles, California, USA May 21-25, 2000 American Society for Microbiology . ISSN: 1060-2011.

DOCUMENT TYPE: Conference

LANGUAGE: English

SUMMARY LANGUAGE: English

L23 ANSWER 19 OF 27 SCISEARCH COPYRIGHT 2002 ISI (R)

ACCESSION NUMBER: 2000:888017 SCISEARCH

THE GENUINE ARTICLE: 364CN

TITLE: Serologic and nucleic acid evidence for *Babesia microti* in Connecticut (CT) blood donors

AUTHOR: Leiby D A (Reprint); Chung A P; Triano L R; Cable R G; Houghton R L; Lodes M J

CORPORATE SOURCE: AMER RED CROSS, FARMINGTON, CT; CORIXA CORP, SEATTLE, WA; AMER RED CROSS, ROCKVILLE, MD

COUNTRY OF AUTHOR: USA

SOURCE: TRANSFUSION, (OCT 2000) Vol. 40, No. 10, Supp. [S], pp. S2-S2.

Publisher: AMER ASSOC BLOOD BANKS, 8101 GLENBROOK RD, BETHESDA, MD 20814-2749.

ISSN: 0041-1132.

DOCUMENT TYPE: Conference; Journal

Searcher : Shears 308-4994

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FILE SEGMENT: LIFE; CLIN
LANGUAGE: English
REFERENCE COUNT: 0

L23 ANSWER 20 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 11
ACCESSION NUMBER: 1999:388301 HCAPLUS
DOCUMENT NUMBER: 131:85409
TITLE: Antigen and gene sequences for diagnosis and
treatment of Babesia **microti** infection
INVENTOR(S): Reed, Steven G.; Lodes, Michael
J.; Houghton, Raymond;
Sleath, Paul R.; Persing, David;
Bruinsma, Elizabeth
PATENT ASSIGNEE(S): Corixa Corporation, USA; Mayo Foundation for
Medical Education and Research
SOURCE: PCT Int. Appl., 126 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 7
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9929869	A1	19990617	WO 1998-US26437	19981211
W: AU, CA, JP, MX, NZ				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 6214971	B1	20010410	US 1997-990571	19971211
AU 9918204	A1	19990628	AU 1999-18204	19981211
US 2001029295	A1	20011011	US 2000-737178	20001213

PRIORITY APPLN. INFO.:

US 1997-990571	A	19971211
US 1996-723142	A2	19961001
US 1997-845258	A2	19970424
WO 1998-US26437	W	19981211
US 1999-286488	A2	19990405
US 2000-528784	A2	20000317
WO 2000-US9136	A2	20000405
US 2000-569098	A2	20000510
US 2000-605724	A2	20000627
US 2000-656688	A2	20000907
US 2000-685436	A2	20001010

AB Compds. and methods for the diagnosis and treatment of Babesia **microti** infection are disclosed. The compds. provided include polypeptides that contain at least one antigenic portion of a B. **microti** antigen and DNA sequences encoding such polypeptides. Nine antigens share some homol., contain a degenerate repeat of six amino acids with 9-22 repeats occurring in each antigen, and bear some similarity to a Plasmodium falciparum merozoite surface antigen gene. A second group of five antigens bear some homol. to each other but do not show homol. to any previously identified sequences. Two synthetic peptides (BABS-1 and BABS-4) were made to the repeat region of isolated B. **microti** antigen BMNI-3. Twelve BMNI-6 homologs were obtained from hamster and human patients. Antigenic epitopes of such antigens are also provided, together with pharmaceutical compns. and vaccines comprising such polypeptides, DNA sequences or antigenic epitopes. Diagnostic kits contg. such polypeptides, DNA

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sequences or antigenic epitopes and a suitable detection reagent may be used for the detection of *B. microti* infection in patients and biol. samples. Antibodies directed against such polypeptides and antigenic epitopes are also provided.

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 21 OF 27 MEDLINE DUPLICATE 12
ACCESSION NUMBER: 1999221786 MEDLINE
DOCUMENT NUMBER: 99221786 PubMed ID: 10203519
TITLE: Detection of enzootic babesiosis in baboons (*Papio cynocephalus*) and phylogenetic evidence supporting synonymy of the genera *Entopolypoides* and *Babesia*.
AUTHOR: Bronsdon M A; Homer M J; Magera J M; Harrison C; Andrews R G; Bielitzki J T; Emerson C L; Persing D H; Fritsche T R
CORPORATE SOURCE: Regional Primate Research Center, University of Washington School of Medicine, Seattle, Washington 98195, USA.
CONTRACT NUMBER: AI35191 (NIAID)
AI41103-01 (NIAID)
RR00166 (NCRR)
SOURCE: JOURNAL OF CLINICAL MICROBIOLOGY, (1999 May) 37 (5) 1548-53.
Journal code: 7505564. ISSN: 0095-1137.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: Priority Journals
OTHER SOURCE: GENBANK-AF081465
ENTRY MONTH: 199905
ENTRY DATE: Entered STN: 19990525
Last Updated on STN: 20000303
Entered Medline: 19990507

AB Blood smear evaluation of two baboons (*Papio cynocephalus*) experiencing acute hemolytic crises following experimental stem cell transplantation revealed numerous intraerythrocytic organisms typical of the genus *Babesia*. Both animals had received whole-blood transfusions from two baboon donors, one of which was subsequently found to display rare trophozoites of *Entopolypoides macaci*. An investigation was then undertaken to determine the prevalence of hematozoa in baboons held in our primate colony and to determine the relationship, if any, between the involved species. Analysis of thick and thin blood films from 65 healthy baboons (23 originating from our breeding facility, 26 originating from an out-of-state breeding facility, and 16 imported from Africa) for hematozoa revealed rare *E. macaci* parasites in 31%, with respective prevalences of 39, 35, and 12%. Phylogenetic analysis of nuclear small-subunit rRNA gene sequences amplified from peripheral blood of a baboon chronically infected with *E. macaci* demonstrated this parasite to be most closely related to *Babesia microti* (97.9% sequence similarity); sera from infected animals did not react in indirect fluorescent-antibody tests with *Babesia microti* antigen, however, suggesting that they represent different species. These results support an emerging view that the genus *Entopolypoides* Mayer 1933 is synonymous with that of the genus *Babesia* Starcovici 1893 and that the morphological variation noted

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among intracellular forms is a function of alteration in host immune status. The presence of an underrecognized, but highly enzootic, *Babesia* sp. in baboons may result in substantial, unanticipated impact on research programs. The similarity of this parasite to the known human pathogen *B. microti* may also pose risks to humans undergoing xenotransplantation, mandating effective screening of donor animals.

L23 ANSWER 22 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 13

ACCESSION NUMBER: 1999:316369 HCAPLUS

DOCUMENT NUMBER: 131:143166

TITLE: A multi-epitope synthetic peptide and recombinant protein for the detection of antibodies to *Trypanosoma cruzi* in radioimmunoprecipitation-confirmed and consensus-positive sera

AUTHOR(S): Houghton, Raymond L.; Benson, Darin R.; Reynolds, Lisa D.; McNeill, Patricia D.; Sleath, Paul R.; Lodes, Michael J.; Skeiky, Yasir A. W.; Leiby, David A.; Badaro, Roberto; Reed, Steven G.

CORPORATE SOURCE: Corixa Corporation, Seattle, WA, 98104, USA
SOURCE: Journal of Infectious Diseases (1999), 179(5), 1226-1234

CODEN: JIDIAQ; ISSN: 0022-1899

PUBLISHER: University of Chicago Press

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Peptide epitopes of *Trypanosoma cruzi* have been identified through expression cloning. A tripeptide (2/D/E) contg. three epitopes (TcD, TcE, PEP-2) was used in ELISA to detect antibodies to *T. cruzi* in 239 of 240 consensus-pos. sera and 41 of 42 sera confirmed pos. by radioimmunopptn. assay. The 1 discrepant consensus-pos. serum was used to expression-clone a novel gene that contained a repeat sequence. A peptide corresponding to this sequence, TcLol.2, was specific for *T. cruzi*. This antigen detected the discrepant consensus-pos. serum and enhanced reactivity of low-pos. sera in the tripeptide assay. A branched synthetic peptide, 2/D/E/Lol.2, or a linear recombinant, r2/D/E/Lol.2, realized all of the diagnostic features of the 4 epitopes, including the ability to boost reactivity of low-reactive sera. Thus, peptides and recombinants contg. multiple repeat epitopes are powerful tools for developing assays for *T. cruzi* antibody detection and have direct application in blood screening.

REFERENCE COUNT: 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 23 OF 27 HCAPLUS COPYRIGHT 2002 ACS DUPLICATE 14

ACCESSION NUMBER: 1998:229027 HCAPLUS

DOCUMENT NUMBER: 128:292989

TITLE: Antigens of *Babesia microtia* and their use in the diagnosis and treatment of infection

INVENTOR(S): Reed, Steven G.; Lodes, Michael J.; Houghton, Raymond; Sleath, Paul R.

PATENT ASSIGNEE(S): Corixa Corp., USA

SOURCE: Eur. Pat. Appl., 113 pp.

Searcher : Shears 308-4994

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CODEN: EPXXDW
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 7
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 834567	A2	19980408	EP 1997-117067	19971001
EP 834567	A3	19990331		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
US 6306396	B1	20011023	US 1996-723142	19961001
US 6183976	B1	20010206	US 1997-845258	19970424
PRIORITY APPLN. INFO.:			US 1996-723142	A 19961001
			US 1997-845258	A 19970424

AB Antigens and epitopes of Babesia **microti** that can be used in the diagnosis and treatment of infection are described. Genes for the antigens or antibodies against them can be used in the detection of B. **microti**. CDNAS for these antigens were cloned by screening an expression library in .lambda.ZAP with antiserum to B. **microti**.

L23 ANSWER 24 OF 27 WPIDS (C) 2002 THOMSON DERWENT
ACCESSION NUMBER: 1998-609891 [51] WPIDS
CROSS REFERENCE: 2000-160675 [11]; 2002-062250 [01]
DOC. NO. CPI: C1998-182724
TITLE: Poly peptide(s) comprising immunogenic portion of Ehrlichia antigen - and encoding DNA sequences, useful for e.g. diagnosis and treatment of Ehrlichia infection, especially human granulocytic ehrlichiosis.
DERWENT CLASS: A96 B04 D16
INVENTOR(S): HOUGHTON, R; LODES, M J;
REED, S G; HOUGHTON, R L
PATENT ASSIGNEE(S): (CORI-N) CORIXA CORP
COUNTRY COUNT: 81
PATENT INFORMATION:

PATENT NO	KIND	DATE	WEEK	LA	PG
WO 9842740	A2	19981001	(199851)*	EN	140
RW: AT BE CH DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW NL OA PT SD SE SZ UG ZW					
W: AL AM AT AU BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG UZ VN YU ZW					
AU 9865794	A	19981020	(199909)		
EP 1007550	A2	20000614	(200033)	EN	
R: AT BE CH DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE					
US 6207169	B1	20010327	(200119)		
US 6231869	B1	20010515	(200129)		
JP 2002515763	W	20020528	(200238)		139

APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
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Searcher : Shears 308-4994

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WO 9842740	A2	WO 1998-US5695	19980323
AU 9865794	A	AU 1998-65794	19980323
EP 1007550	A2	EP 1998-911966	19980323
		WO 1998-US5695	19980323
US 6207169	B1 CIP of	US 1997-821324	19970321
		US 1997-975762	19971120
US 6231869	B1	US 1997-821324	19970321
JP 2002515763 W		JP 1998-545891	19980323
		WO 1998-US5695	19980323

FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 9865794	A Based on	WO 9842740
EP 1007550	A2 Based on	WO 9842740
JP 2002515763 W	Based on	WO 9842740

PRIORITY APPLN. INFO: US 1997-975762 19971120; US 1997-821324
19970321

AN 1998-609891 [51] WPIDS
CR 2000-160675 [11]; 2002-062250 [01]
AB WO 9842740 A UPAB: 20020701

A polypeptide comprising an immunogenic portion of an Ehrlichia antigen (or variant differing by conservative substitutions and/or modifications) is new, in which the antigen has amino acid sequence encoded by:

- (i) one of nineteen 201-7091 bp sequences given in the specification, encoding Ehrlichia antigens;
- (ii) sequences hybridising to (i); or
- (iii) complements of (i) or (ii).

Also claimed are:

(1) an antigenic epitope of an Ehrlichia antigen with the 41 or 125 amino acid sequences (I) or (II) given in the specification, and a second polypeptide comprising at least two such antigenic epitopes.

(2) DNA molecules encoding polypeptides as above;

(3) expression vectors comprising (2);

(4) host cells (e.g. E. coli, yeast or mammalian) transformed with (3);

(5) fusion proteins comprising at least one of polypeptides and/or at least one of antigenic epitopes;

(6) diagnostic kits comprising at least one polypeptide/antigenic epitope/fusion protein (optionally immobilised on solid support e.g. nitrocellulose) plus detection reagent (optionally comprising reporter group e.g. radioisotope conjugated to binding agent e.g. anti-immunoglobulin; and

(7) monoclonal/polyclonal antibodies binding to polypeptides/antigenic epitope.

USE - The polypeptides are useful in the treatment of Ehrlichia infection, and as vaccines for the prevention of infection (claimed). They preferably comprise an immunogenic portion of an Ehrlichia antigen associated with human granulocytic ehrlichiosis (or a variant) (claimed) and are thus especially useful in the treatment of human granulocytic ehrlichiosis (HGE). The polypeptides, antigenic epitopes or DNA molecules can be combined with a suitable carrier in pharmaceutical compositions (claimed).

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Such compositions and the vaccines of (7) are useful to manufacture medicaments for inducing protective immunity against Ehrlichia infection in patients (claimed), especially against HGE. The polypeptides/antigenic epitopes/fusion proteins are also useful to detect such infections in patients, by contacting biological samples (e.g. serum, and especially whole blood (claimed) with at least 1 polypeptide/antigenic epitope/fusion protein and detecting antibody binding (claimed). They can also be used to produce antibodies useful in diagnosis of such infections. The DNA molecules of (2) are similarly useful for diagnosing such infections (claimed). HGE is caused by a rodent bacterium normally transmitted to humans by the same tick which transmits Lyme disease and babesiosis. Co-infection with these diseases is thus possible, and the polypeptides/antigenic epitopes/fusion proteins may be used in methods to detect at least one of Ehrlichia infection, Lyme disease or *B. microti* infection in patients, by contacting samples (e.g. whole blood, saliva etc.) with at least one polypeptide/antigenic epitope/fusion protein, a Lyme disease antigen and a *B. microti* antigen, and detecting antibody binding (claimed); kits are provided (claimed).

Dwg.0/2

L23 ANSWER 25 OF 27 CONFSCI COPYRIGHT 2002 CSA

ACCESSION NUMBER: 1998:37806 CONFSCI

DOCUMENT NUMBER: 98-037806

TITLE: Immunoreactivity of recombinant antigens of Babesia *microti* isolated using serological expression cloning

AUTHOR: Houghton, R.L.; Bruinsma, E.S.; Moro, M.H.; Krause, P.J.; Reynolds, L.D.; Mohamath, R.; Benson, D.R.; Lodes, M.J.; Reed, S.G.; Persing, D.H.

CORPORATE SOURCE: Corixa Corporation, Seattle, WA, USA

SOURCE: ASTMH, 60 Revere Drive, Suite 500, Northbrook, IL 60062, USA, Abstracts available. Price \$10. Poster Paper No. 558.
Meeting Info.: 981 5000: 46th Annual Meeting of the American Society of Tropical Medicine and Hygiene (9815000). Lake Buena Vista, FL (USA). 7-11 Dec 1997. American Society of Tropical Medicine and Hygiene.

DOCUMENT TYPE: Conference

FILE SEGMENT: DCCP

LANGUAGE: English

L23 ANSWER 26 OF 27 CONFSCI COPYRIGHT 2002 CSA

ACCESSION NUMBER: 2002:14683 CONFSCI

DOCUMENT NUMBER: 02-014683

TITLE: Identification and partial characterization of secreted antigens from Babesia *microti* using a novel approach

AUTHOR: Homer, M.J.; Lodes, M.J.; Reynolds, L.D.; Houghton, R.L.; Persing, D.H.

CORPORATE SOURCE: Corixa Corporation, Seattle, WA, USA

SOURCE: American Society for Tropical Medicine, 60 Revere Dr., Suite 500, Northbrook, IL 60062, USA; phone: 847-480-9592; fax: 847-480-9282; email: astmh@astmh.org; URL: www.astmh.org. Poster Paper No.

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09/853079

671.

Meeting Info.: 000 5775: 50th Annual Meeting of the American Society for Tropical Medicine (0005775). Atlanta, GA (USA). 11-15 Nov 2001. Bill and Melinda Gates Foundation, Glaxo SmithKline, Oravax Inc., Berna Products.

DOCUMENT TYPE: Conference
FILE SEGMENT: DCCP
LANGUAGE: English

L23 ANSWER 27 OF 27 CONFSCI COPYRIGHT 2002 CSA

ACCESSION NUMBER: 1998:37803 CONFSCI

DOCUMENT NUMBER: 98-037803

TITLE: Geographic variation within a gene encoding an immunoreactive protein from *Babesia microti*

AUTHOR: Bruinsma, E.S.; Lodes, M.J.; Moro, M.; Krause, P.J.; Reynolds, L.D.; Mohamath, R.; Benson, D.R.; Houghton, R.L.; Reed, S.G.; Persing, D.H.

CORPORATE SOURCE: Dep. Med. and Pathol., Mayo Clinic, Rochester, MN, USA

SOURCE: ASTMH, 60 Revere Drive, Suite 500, Northbrook, IL 60062, USA, Abstracts available. Price \$10. Poster Paper No. 554.

Meeting Info.: 981 5000: 46th Annual Meeting of the American Society of Tropical Medicine and Hygiene (9815000). Lake Buena Vista, FL (USA). 7-11 Dec 1997. American Society of Tropical Medicine and Hygiene.

DOCUMENT TYPE: Conference
FILE SEGMENT: DCCP
LANGUAGE: English

FILE 'HOME' ENTERED AT 12:24:46 ON 30 AUG 2002

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claim 4

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: August 30, 2002, 13:18:44 ; Search time 51.8 Seconds
(without alignments)
12.866 Million cell updates/sec

Title: BASK-853-CLAIM4

Perfect score: 21

Sequence: 1 eagrxx 6

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 747574 seqs, 111073796 residues

Total number of hits satisfying chosen parameters: 747574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 50 summaries

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- 4: /SIDS1/gcgdata/hold-geneseq/geneq-emb1/AA1983.DAT.*
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- 9: /SIDS1/gcgdata/hold-geneseq/geneq-emb1/AA1988.DAT.*
- 10: /SIDS1/gcgdata/hold-geneseq/geneq-emb1/AA1989.DAT.*
- 11: /SIDS1/gcgdata/hold-geneseq/geneq-emb1/AA1990.DAT.*
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- 20: /SIDS1/gcgdata/hold-geneseq/geneq-emb1/AA1999.DAT.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19	90.5	47	19 AAW77585	Staphylococcus aur
2	19	90.5	81	22 AAU61772	Propionibacterium
3	19	90.5	97	22 AAU41551	Propionibacterium
4	19	90.5	100	22 ABG27830	Novel human diago
5	19	90.5	113	22 AAU43242	Propionibacterium
6	19	90.5	117	22 AAM84257	Human immune/haema
7	19	90.5	118	20 AAW59973	Human endometrium
8	19	90.5	159	19 AAW31552	Collagen binding p
9	19	90.5	168	22 AAEL1855	Staphylococcus aur
10	19	90.5	177	14 AAR39711	A. oryzae wt neutr
11	19	90.5	177	14 AAR39712	A. oryzae C6A neut

12	19	90.5	177	14	AAAR39713	A. oryzae C78A neu
13	19	90.5	207	21	AAAB24437	Human secreted pro
14	19	90.5	211	19	AAAB31553	Collagen binding p
15	19	90.5	225	22	ABG03051	Novel human diago
16	19	90.5	229	21	AAAG05085	Arabidopsis thalia
17	19	90.5	242	22	AAAM25472	Human protein sequ
18	19	90.5	243	21	AAAG05084	Arabidopsis thalia
19	19	90.5	243	21	AAAY5431	Human calcium chan
20	19	90.5	257	21	AAAG05083	Arabidopsis thalia
21	19	90.5	264	22	AAU68590	Human novel cytoki
22	19	90.5	270	22	ABG69721	Drosophila melanog
23	19	90.5	270	22	ABG03786	Novel human diago
24	19	90.5	273	22	AAU31144	Novel human secret
25	19	90.5	290	22	AAU47789	Propionibacterium
26	19	90.5	305	22	ABG02081	Novel human diago
27	19	90.5	323	22	AAAB9464	Human protein sequ
28	19	90.5	330	11	AAAR05528	High density lipop
29	19	90.5	332	21	AAAY5898	Human myristoylate
30	19	90.5	332	21	AAAY5899	Human myristoylate
31	19	90.5	333	21	AAAY97409	zebrafish Hsp-bind
32	19	90.5	341	22	AAAG90789	C glutamicum prote
33	19	90.5	341	22	AAAB79110	Corynebacterium gl
34	19	90.5	341	22	AAAB79144	Corynebacterium gl
35	19	90.5	349	19	AAAW44368	Aspergillus nidula
36	19	90.5	352	12	AAAR14147	Pre-pro neutral pr
37	19	90.5	363	22	AAU36356	Pseudomonas aerugi
38	19	90.5	366	20	AAAY05663	Maize caffeic O-me
39	19	90.5	377	21	AAAG20561	Arabidopsis thalia
40	19	90.5	377	21	AAAG41800	Arabidopsis thalia
41	19	90.5	386	21	AAAG24042	Arabidopsis thalia
42	19	90.5	390	8	AAAP70581	Protease biosynthe
43	19	90.5	393	20	AAAY35147	Chlamydia pneumoni
44	19	90.5	422	20	AAAY28643	Human serine prote
45	19	90.5	422	20	AAAB74691	Human protease and
46	19	90.5	427	21	AAAG41388	Arabidopsis thalia
47	19	90.5	431	21	AAAG41387	Arabidopsis thalia
48	19	90.5	435	21	AAAG24041	Arabidopsis thalia
49	19	90.5	440	21	AAAG24040	Arabidopsis thalia
50	19	90.5	441	14	AAAR31955	Sequence encoded b

ALIGNMENTS

RESULT 1
AAW77585
ID AAW77585 standard; Protein; 47 AA.
XX
AC AAW77585;
XX
DT 30-OCT-1998 (first entry)
XX
DE Staphylococcus aureus protein of unknown function.
XX
KW Staphylococcus aureus protein; immune response induction; eye infection;
KW antibody production; T-cell immune response; gastrointestinal infection;
KW respiratory infection; inhibitor; bacterial infection; cardiac infection;
KW central nervous system; kidney infection; urinary tract infection;
KW antimicrobial compound identification; broad spectrum antibiotic;
KW therapy.

OS	Staphylococcus aureus.
XX	
XX	Key Location/Qualifiers
PH	Misc-difference 1..47
FT	/note= "residues designated X are unspecified, and represented as xaa in the specification"
XX	
XX	EP841394-A2.
XX	
XX	13-MAY-1998.
PD	
XX	24-SEP-1997; 97EP-0307485.

```

XX PR 24-SEP-1996; 96US-0027032.
XX (SMIK ) SMITHKLINE BEECHAM CORP.
XX PA (SMIK ) SMITHKLINE BEECHAM PLC.
XX XX
XX Black MT, Burnham MKR, Hodgson JE, Knowles DJC;
PI Lonetto MA, Nicholas RO, Pratt JM, Reichard RW, Rosenberg M;
PI Ward JM;
XX XX
XX WPI: 1998-252940/23.
DR N-PSDB; AAV53383.
XX XX
XX New nucleic acid sequences from Staphylococcus aureus WCHU29 -
PT useful in vaccines and for treatment of bacterial infections of e.g.
PT respiratory tract and central nervous system
XX XX
XX Claim 11; Page 267; 390pp; English.
XX XX
XX This sequence represents a Staphylococcus aureus protein of unknown
CC function, and is encoded by a DNA sequence of the invention.
CC The DNA sequences were isolated from Staphylococcus aureus WCHU29
CC (NCIMB 40771). Host cells containing the DNA sequences are used to
CC produce polypeptides or fragments. The proteins are used in the treatment
CC of disease, for inducing an immune response by administering them, to
CC produce antibody and/or T-cell immune response. Antagonists of the
CC proteins are used for the inhibition of bacterial polypeptides.
CC Conditions which may be treated include bacterial infections, especially
CC respiratory, cardiac, gastrointestinal, central nervous, eye, kidney,
CC urinary tract, skin, bones and joints. The proteins can also be used to
CC identify antimicrobial compounds which are broad spectrum antibiotics,
CC especially useful in the treatment of H. pylori infection.
XX XX
XX Sequence 47 AA;

Query Match 90.5%; Score 19; DB 19; Length 47;
Best Local Similarity 66.7%; Pred. No. 2.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
Db 31 eagats 36
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RESULT 2
AAU61772
XX AAU61772 standard; Protein; 81 AA.
XX AC AAU61772;
XX DT 27-FEB-2002 (first entry)
XX DE Propionibacterium acnes immunogenic protein #22668.
XX KW SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;
KW dermatological; osteopathic; neuroprotectant.
XX OS Propionibacterium acnes.
XX PN WO200181581-A2.
XX PD 01-NOV-2001.
XX PF 20-APR-2001; 2001WO-US12865.
XX PR 21-APR-2000; 2000US-199047P.
XX PR 02-JUN-2000; 2000US-208841P.
XX PR 07-JUL-2000; 2000US-216747P.
XX PA (CORI-) CORIXA CORP.

XX Skelky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;
PI L'maisonneuve J, Zhang Y, Jen S, Carter D;
XX XX
XX WPI: 2001-616774/71.
DR N-PSDB; AAS59620.
XX XX
XX Propionibacterium acnes polypeptides and nucleic acids useful for
PT vaccinating against and diagnosing infections, especially useful for
PT treating acne vulgaris -
XX XX
XX Example 1; SEQ ID No 22967; 1069pp; English.
XX XX
XX Sequences AAU39105-AAU68017 represent Propionibacterium acnes immunogenic
CC polypeptides. The proteins and their associated DNA sequences are used in
CC the treatment, prevention and diagnosis of medical conditions caused by
CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,
CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.
CC P. acnes is also involved in infections of bone, joints and the central
CC nervous system, however it is particularly involved in the inflammatory
CC lesions associated with acne vulgaris. A method for detecting the
CC presence or absence of P. acnes in a patient comprises contacting a
CC sample with a binding agent that binds to the proteins of the invention
CC and determining the amount of bound protein in the sample. The
CC polypeptides may be used as antigens in the production of antibodies
CC specific for P. acnes proteins. These antibodies can be used to
CC downregulate expression and activity of P. acnes polypeptides and
CC therefore treat P. acnes infections. The antibodies may also be used as
CC diagnostic agents for determining P. acnes presence, for example, by
CC enzyme linked immunosorbent assay (ELISA).
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX XX
XX Sequence 81 AA;

Query Match 90.5%; Score 19; DB 22; Length 81;
Best Local Similarity 66.7%; Pred. No. 5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
Db 20 eagats 25
||| |

RESULT 3
AAU41551
XX AAU41551 standard; Protein; 97 AA.
XX AC AAU41551;
XX DT 13-FEB-2002 (first entry)
XX DE Propionibacterium acnes immunogenic protein #2447.
XX KW SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;
KW dermatological; osteopathic; neuroprotectant.
XX OS Propionibacterium acnes.
XX PN WO200181581-A2.
XX PD 01-NOV-2001.
XX PF 20-APR-2001; 2001WO-US12865.
XX PR 21-APR-2000; 2000US-199047P.
XX PR 02-JUN-2000; 2000US-208841P.
XX PR 07-JUL-2000; 2000US-216747P.
XX XX

```

PA (CORI-) CORIXA CORP.
 XX Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;
 PI L'maisonneuve J, Zhang Y, Jen S, Carter D;
 XX
 DR WPI: 2001-616774/71.
 DR N-PSDB; AAS59515.
 XX
 PT Propionibacterium acnes polypeptides and nucleic acids useful for
 PT vaccinating against and diagnosing infections, especially useful for
 PT treating acne vulgaris -
 XX
 XX Example 1; SEQ ID No 2746; 1069pp; English.
 PS
 XX Sequences AAU39105-AAU68017 represent Propionibacterium acnes immunogenic
 CC polypeptides. The proteins and their associated DNA sequences are used in
 CC the treatment, prevention and diagnosis of medical conditions caused by
 CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,
 CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.
 CC P. acnes is also involved in infections of bone, joints and the central
 CC nervous system, however it is particularly involved in the inflammatory
 CC lesions associated with acne vulgaris. A method for detecting the
 CC presence or absence of P. acnes in a patient comprises contacting a
 CC sample with a binding agent that binds to the proteins of the invention
 CC and determining the amount of bound protein in the sample. The
 CC polypeptides may be used as antigens in the production of antibodies
 CC specific for P. acnes proteins. These antibodies can be used to
 CC downregulate expression and activity of P. acnes polypeptides and
 CC therefore treat P. acnes infections. The antibodies may also be used as
 CC diagnostic agents for determining P. acnes presence, for example, by
 CC enzyme linked immunosorbent assay (ELISA).
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 97 AA;

Query Match 90.5%; Score 19; DB 22; Length 97;
 Best Local Similarity 66.7%; Pred. No. 6e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 29 eagsas 34

RESULT 4
 ABG27830
 ID ABG27830 standard; Protein; 100 AA.

XX AC ABG27830;

XX DT 18-FEB-2002 (first entry)

XX DE Novel human diagnostic protein #27821.

XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 XX KW food supplement; medical imaging; diagnostic; genetic disorder.

XX OS Homo sapiens.

XX PN WO200175067-A2.

XX PD 11-OCT-2001.

XX PF 30-MAR-2001; 2001WO-US08631.

XX PR 31-MAR-2000; 2000US-0540217.

XX PR 23-AUG-2000; 2000US-0649167.

XX PA (HYSE-) HYSEQ INC.

XX

PI Drmanac RT, Liu C, Tang YT;
 XX
 XX WPI: 2001-639362/73.
 DR N-PSDB; AAS92017.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 XX Claim 20; SEQ ID No 58189; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human
 CC diagnostic amino acid sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX

SQ Sequence 100 AA;

Query Match 90.5%; Score 19; DB 22; Length 100;
 Best Local Similarity 66.7%; Pred. No. 6.1e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 95 eagats 100

RESULT 5
 AAU43242

ID AAU43242 standard; Protein; 113 AA.

XX AC AAU43242;

XX DT 27-FEB-2002 (first entry)

XX DE Propionibacterium acnes immunogenic protein #4138.

XX KW SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
 XX KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;
 XX KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;
 XX KW dermatological; osteopathic; neuroprotectant.

XX OS Propionibacterium acnes.

XX PN WO200181581-A2.

XX PD 01-NOV-2001.

XX PF 20-APR-2001; 2001WO-US12865.

XX PR 21-APR-2000; 2000US-199047P.

XX PR 02-JUN-2000; 2000US-208841P.

XX PR 07-JUL-2000; 2000US-216747P.

XX

PA (CORI-) CORIXA CORP.
XX Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;
PI L'maisonneuve J, Zhang Y, Jen S, Carter D;
XX WPI; 2001-616774/71.
DR N-PSDB; AAS95920.
XX
PT Propionibacterium acnes polypeptides and nucleic acids useful for
PT vaccinating against and diagnosing infections, especially useful for
PT treating acne vulgaris -
XX
PS Example 1; SEQ ID No 4437; 1069pp; English.
XX
CC Sequences AAU39105-AAU68017 represent Propionibacterium acnes immunogenic
CC polypeptides. The proteins and their associated DNA sequences are used in
CC the treatment, prevention and diagnosis of medical conditions caused by
CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,
CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.
CC P. acnes is also involved in infections of bone, joints and the central
CC nervous system, however it is particularly involved in the inflammatory
CC lesions associated with acne vulgaris. A method for detecting the
CC presence or absence of P. acnes in a patient comprises contacting a
CC sample with a binding agent that binds to the proteins of the invention
CC and determining the amount of bound protein in the sample. The
CC polypeptides may be used as antigens in the production of antibodies
CC specific for P. acnes proteins. These antibodies can be used to
CC downregulate expression and activity of P. acnes polypeptides and
CC therefore treat P. acnes infections. The antibodies may also be used as
CC diagnostic agents for determining P. acnes presence, for example, by
CC enzyme linked immunosorbent assay (ELISA).
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
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SQ Sequence 113 AA;

Query Match 90.5%; Score 19; DB 22; Length 113;
Best Local Similarity 66.7%; Pred. No. 6.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
Db 99 eagtas 104

RESULT 6
AAM84257
ID AAM84257 standard; Protein: 117 AA.
AC AAM84257;
XX
XX 07-NOV-2001 (first entry)
XX
XX Human immune/haematopoietic antigen SEQ ID NO:11850.
XX
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
KW cytostatic; gene therapy; vaccine; metastasis.
KW Homo sapiens.
OS
XX WO200157182-A2.
PN
XX 09-AUG-2001.
PD
XX 17-JAN-2001; 2001WO-US01354.
PF
XX 31-JAN-2000; 2000US-0179065.
PR 04-FEB-2000; 2000US-0180628.
PR 24-FEB-2000; 2000US-0184664.
PR 02-MAR-2000; 2000US-0186350.
PR 16-MAR-2000; 2000US-0189874.
PR
17-MAR-2000; 2000US-0190076.
PR 18-APR-2000; 2000US-0198123.
PR 19-MAY-2000; 2000US-0205515.
PR 07-JUN-2000; 2000US-0209467.
PR 28-JUN-2000; 2000US-0214886.
PR 30-JUN-2000; 2000US-0215135.
PR 07-JUL-2000; 2000US-0216647.
PR 07-JUL-2000; 2000US-0216880.
PR 11-JUL-2000; 2000US-0217487.
PR 11-JUL-2000; 2000US-0217496.
PR 14-JUL-2000; 2000US-0218290.
PR 26-JUL-2000; 2000US-0220963.
PR 26-JUL-2000; 2000US-0220964.
PR 14-AUG-2000; 2000US-0224518.
PR 14-AUG-2000; 2000US-0224519.
PR 14-AUG-2000; 2000US-0225213.
PR 14-AUG-2000; 2000US-0225214.
PR 14-AUG-2000; 2000US-0225266.
PR 14-AUG-2000; 2000US-0225267.
PR 14-AUG-2000; 2000US-0225268.
PR 14-AUG-2000; 2000US-0225270.
PR 14-AUG-2000; 2000US-0225447.
PR 14-AUG-2000; 2000US-0225757.
PR 14-AUG-2000; 2000US-0225758.
PR 14-AUG-2000; 2000US-0225759.
PR 18-AUG-2000; 2000US-0226279.
PR 22-AUG-2000; 2000US-0226681.
PR 22-AUG-2000; 2000US-0226868.
PR 22-AUG-2000; 2000US-0227182.
PR 23-AUG-2000; 2000US-0227009.
PR 30-AUG-2000; 2000US-0228924.
PR 01-SEP-2000; 2000US-0229287.
PR 01-SEP-2000; 2000US-0229343.
PR 01-SEP-2000; 2000US-0229344.
PR 01-SEP-2000; 2000US-0229345.
PR 05-SEP-2000; 2000US-0229509.
PR 05-SEP-2000; 2000US-0229513.
PR 06-SEP-2000; 2000US-0230437.
PR 06-SEP-2000; 2000US-0230438.
PR 08-SEP-2000; 2000US-0231242.
PR 08-SEP-2000; 2000US-0231243.
PR 08-SEP-2000; 2000US-0231244.
PR 08-SEP-2000; 2000US-0231413.
PR 08-SEP-2000; 2000US-0231414.
PR 08-SEP-2000; 2000US-0232080.
PR 08-SEP-2000; 2000US-0232081.
PR 12-SEP-2000; 2000US-0231968.
PR 14-SEP-2000; 2000US-0232397.
PR 14-SEP-2000; 2000US-0232398.
PR 14-SEP-2000; 2000US-0232399.
PR 14-SEP-2000; 2000US-0232400.
PR 14-SEP-2000; 2000US-0232401.
PR 14-SEP-2000; 2000US-0233063.
PR 14-SEP-2000; 2000US-0233064.
PR 14-SEP-2000; 2000US-0233065.
PR 21-SEP-2000; 2000US-0234223.
PR 21-SEP-2000; 2000US-0234274.
PR 25-SEP-2000; 2000US-0234997.
PR 25-SEP-2000; 2000US-0234998.
PR 26-SEP-2000; 2000US-0235484.
PR 27-SEP-2000; 2000US-0235834.
PR 27-SEP-2000; 2000US-0235836.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236367.
PR 29-SEP-2000; 2000US-0236368.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237040.
PR 13-OCT-2000; 2000US-0239935.

PR 13-OCT-2000; 2000US-0239937.
 PR 20-OCT-2000; 2000US-0240360.
 PR 20-OCT-2000; 2000US-0241221.
 PR 20-OCT-2000; 2000US-0241785.
 PR 20-OCT-2000; 2000US-0241786.
 PR 20-OCT-2000; 2000US-0241787.
 PR 20-OCT-2000; 2000US-0241808.
 PR 20-OCT-2000; 2000US-0241809.
 PR 20-OCT-2000; 2000US-0241826.
 PR 01-NOV-2000; 2000US-0241617.
 PR 08-NOV-2000; 2000US-0246474.
 PR 08-NOV-2000; 2000US-0246475.
 PR 08-NOV-2000; 2000US-0246476.
 PR 08-NOV-2000; 2000US-0246477.
 PR 08-NOV-2000; 2000US-0246478.
 PR 08-NOV-2000; 2000US-0246523.
 PR 08-NOV-2000; 2000US-0246524.
 PR 08-NOV-2000; 2000US-0246525.
 PR 08-NOV-2000; 2000US-0246526.
 PR 08-NOV-2000; 2000US-0246527.
 PR 08-NOV-2000; 2000US-0246528.
 PR 08-NOV-2000; 2000US-0246532.
 PR 08-NOV-2000; 2000US-0246609.
 PR 08-NOV-2000; 2000US-0246610.
 PR 08-NOV-2000; 2000US-0246611.
 PR 17-NOV-2000; 2000US-0246613.
 PR 17-NOV-2000; 2000US-0249207.
 PR 17-NOV-2000; 2000US-0249208.
 PR 17-NOV-2000; 2000US-0249209.
 PR 17-NOV-2000; 2000US-0249210.
 PR 17-NOV-2000; 2000US-0249211.
 PR 17-NOV-2000; 2000US-0249212.
 PR 17-NOV-2000; 2000US-0249213.
 PR 17-NOV-2000; 2000US-0249214.
 PR 17-NOV-2000; 2000US-0249215.
 PR 17-NOV-2000; 2000US-0249216.
 PR 17-NOV-2000; 2000US-0249217.
 PR 17-NOV-2000; 2000US-0249218.
 PR 17-NOV-2000; 2000US-0249219.
 PR 17-NOV-2000; 2000US-0249244.
 PR 17-NOV-2000; 2000US-0249245.
 PR 17-NOV-2000; 2000US-0249264.
 PR 17-NOV-2000; 2000US-0249265.
 PR 17-NOV-2000; 2000US-0249297.
 PR 17-NOV-2000; 2000US-0249299.
 PR 17-NOV-2000; 2000US-0249300.
 PR 01-DEC-2000; 2000US-0250160.
 PR 01-DEC-2000; 2000US-0250391.
 PR 05-DEC-2000; 2000US-0251030.
 PR 05-DEC-2000; 2000US-0251988.
 PR 06-DEC-2000; 2000US-0256719.
 PR 06-DEC-2000; 2000US-0251479.
 PR 08-DEC-2000; 2000US-0251856.
 PR 08-DEC-2000; 2000US-0251868.
 PR 08-DEC-2000; 2000US-0251869.
 PR 08-DEC-2000; 2000US-0251989.
 PR 11-DEC-2000; 2000US-0251990.
 PR 05-JAN-2001; 2000US-0254097.
 PR 05-JAN-2001; 2000US-0259678.
 (HUMA-) HUMAN GENOME SCI INC.
 PA Rosen CA, Barash SC, Ruben SM;
 XX N-PSDB; AAK57038.
 DR WPI; 2001-483426/52.
 DR N-PSDB; AAK57038.
 XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
 PT useful for preventing, diagnosing and/or treating cancers and
 PT metastasis -
 PT
 XX Claim 11; SEQ ID NO 11850; 3071pp + Sequence Listing; English.
 PS AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)
 XX
 CC

CC amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic
 CC activity, and can be used in gene therapy and vaccine production. (I)
 CC proteins and polynucleotides may be used in the prevention, diagnosis and
 CC treatment of diseases associated with inappropriate (I) expression. For
 CC example, they may be used to treat disorders associated with decreased
 CC expression by rectifying mutations or deletions in a patient's genome
 CC that affect the activity of (I) by expressing inactive proteins or to
 CC supplement the patients own production of (I). Additionally, (I)
 CC polynucleotides may be used to produce the secreted (I), by inserting
 CC the nucleic acids into a host cell and culturing the cell to express the
 CC protein. (I) proteins and polynucleotides may be used to prevent,
 CC diagnose and treat immune/hematopoietic-related diseases, especially
 CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
 CC to AAK87694 represent human immune/haematopoietic antigen genomic
 CC sequences from the present invention. AAK54942 to AAK54950 and AAM82169
 CC represent sequences used in the exemplification of the present invention.
 XX

SQ Sequence 117 AA;

Query Match 90.5%; Score 19; DB 22; Length 117;
 Best Local Similarity 66.7%; Pred. No. 7.2e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 78 eagtas 83

RESULT 7

AAV59973
 ID AAV59973 standard; Protein; 118 AA.

AC AAV59973;

XX 31-JAN-2000 (first entry)

XX Human endometrium tumour EST encoded protein 33.

XX Endometrium; human; tumour; cancer; anticancer; cytostatic; EST:
 treatment; uterine; gene therapy; expressed sequence tag.

OS Homo sapiens.

XX DEL9817948-A1.

PN 21-OCT-1999.

PD 17-APR-1998; 98DE-1017948.

PF 17-APR-1998; 98DE-1017948.

XX (META-) METAGEN GES GENOMFORSCHUNG MBH.

XX Rosenthal A, Specht T, Hinzmann B, Schmitt A, Pilarsky C, Dahl E;

WPI; 1999-591957/51.

DR N-PSDB; AAZ41991.

XX New nucleic acid sequences expressed in uterine cancer tissues, and
 PT derived polypeptides, for treatment of uterine and endometrial cancer
 PT and identification of therapeutic agents -
 XX Claim 23; Page 288; 444pp; German.

PS This invention describes novel human nucleic acid (cDNA) sequences (A),
 XX that are highly expressed in uterine tumour tissue and which have
 CC anticancer and cytostatic activity. (A) are used (i) for recombinant
 CC expression of polypeptides (B) and (ii) to isolate complete genes. (B)
 CC are used (i) to identify agents suitable for treatment of uterine or
 CC endometrial cancer; (ii) directly for treating these forms of cancer
 CC (including expression from gene therapy vectors) and (iii) for
 CC generation of specific antibodies. (A) are identified by assembling ESTs

CC (expressed sequence tags) from a particular tissue type before comparison
 CC of expression patterns. This allows a significantly longer fragment of
 CC the gene to be revealed, so should reduce the number of failures
 CC associated with the fact that ESTs from different libraries may represent
 CC different parts of the same unknown gene, distorting the estimated
 CC frequency of occurrence in a particular tissue. AAY59941-Y60328 represent
 CC protein fragments encoded by the human endometrium tumour cDNA library
 CC derived EST fragments represented in AA241981-242121.
 XX
 SQ Sequence 118 AA;

Query Match 90.5%; Score 19; DB 20; Length 118;
 Best Local Similarity 66.7%; Pred. No. 7.2e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 34 eagsas 39

RESULT 8
 AAW31552
 ID AAW31552 standard; Protein; 159 AA.
 XX
 AC AAW31552;

DT 21-MAY-1998 (first entry)
 XX
 DE Collagen binding protein M17 epitope.
 XX

KW Collagen binding protein; cna gene; sepsis; infection;
 KW microbial surface component recognising adhesive matrix molecule;
 KW MSCRAMM; adhesin; vaccine; immunisation; diagnosis; therapy;
 KW epitope M17.
 XX

OS Staphylococcus aureus.
 XX

FH Key Location/Qualifiers
 FT Peptide 1..12
 FT /note= "vector pQE30-derived peptide"
 FT Protein 13..159
 FT /note= "epitope M17"
 FT
 XX

PN W09743314-A2.
 XX

PD 20-NOV-1997.
 XX

PE 14-MAY-1997; 97WO-US08210.
 XX

PR 16-MAY-1996; 96US-0017678.
 XX

PA (UABR-) UAB RES FOUND.
 XX

PA (TEXA) UNIV TEXAS A & M SYSTEM.
 XX

PI Hook M, House-Pompeo K, Patti JM, Sthanam N, Symersky J;
 XX

DR WPI; 1998-008801/01.
 XX

DR N-PSDB; AAT93436.
 XX

PT Antibody that interacts with collagen binding domain of
 FT Staphylococcal cna gene product - useful to prevent bacterial sepsis
 FT in animal infected with Staphylococcus aureus
 XX

PS Claim 31; Page 114; 143pp; English.
 XX

CC This protein comprises Staphylococcus aureus collagen binding
 CC protein (CBP) epitope M17, i.e. amino acids 151-297 of full-length
 CC CBP, plus a vector-derived N-terminal peptide. Claimed 441, 849
 CC and 1500 bp nucleic acid sequences (see AAT93436-38) respectively
 CC encode CBP epitopes M17, M31 and M55 (see AAW31552-54) that confer
 CC protection against S. aureus infection. These nucleic acid
 CC sequences can be used in the recombinant production of the CBP

CC epitopes. The CBP protein and antigenic epitopes are contemplated
 CC for use in the treatment of pathological infections, especially to
 CC prevent bacterial adhesion to collagen. The claimed nucleic acids
 CC as well as claimed anti-CBP antibodies will also be of use in
 CC screening, diagnostic and therapeutic applications including active
 CC and passive immunisation and methods for the prevention of
 CC bacterial colonisation in an animal such as a human. The CBP
 CC epitopes are also contemplated for use in the preparation of
 CC vaccines and as carrier proteins in vaccine formulations, as well
 CC as in the formulation of compositions for the prevention of S.
 CC aureus infection.
 XX

SQ Sequence 159 AA;

Query Match 90.5%; Score 19; DB 19; Length 159;
 Best Local Similarity 66.7%; Pred. No. 9.7e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 28 eagtss 33

RESULT 9

AAEI1855

ID AAEI1855 standard; Protein; 168 AA.

AC AAEI1855;

DT 18-DEC-2001 (first entry)

DE Staphylococcus aureus CNA19 protein.

KW Collagen-binding region; CNA19; Staphylococcus aureus infection;

KW Staphylococcus epidermidis infection; vaccine; CNA protein;

KW cross-reactive antibody.
 XX

OS Staphylococcus aureus.
 XX

FH Key Location/Qualifiers

FT Region 24...29
 FT /note= "Beta strand a; this region forms a part of
 FT the trench in the beta sheet"
 FT Region 37...44

FT /note= "Beta strand b; this region forms a part of
 FT the trench in the beta sheet"
 FT Region 55...61

FT /note= "Beta strand c"
 FT Region 65...78

FT /note= "Beta strand d; a portion of this region forms
 FT a part of the trench in the beta sheet"
 FT Region 82...84

FT /note= "Beta strand e; a portion of this region forms
 FT a part of the trench in the beta sheet"
 FT Region 89...92

FT /note= "Alpha helix 1"
 FT Region 93...96

FT /note= "Alpha helix 2"
 FT Region 101...105

FT /note= "Beta strand f"
 FT Region 110...115

FT /note= "Beta strand g"
 FT Region 123...133

FT /note= "Beta strand h; a portion of this region forms
 FT a part of the trench in the beta sheet"
 FT Region 140...149

FT /note= "Beta strand i"
 FT Region 157...167

FT /note= "Beta strand j"
 XX

PN W0200170267-A1.
 XX

PD XX 27-SEP-2001.
 XX PF 19-MAR-2001; 2001WO-US08554.
 XX PR 17-MAR-2000; 2000US-189968P.
 XX PR 25-APR-2000; 2000US-199370P.
 XX PR 15-AUG-2000; 2000US-225402P.
 XX (INH1-) INHIBITEX INC.
 PA (TEXA) UNIV TEXAS A & M SYSTEM.
 PA (UYPA-) UNIV PAVIA.
 XX Hook M, Xu Y, Speziale P, Visai L, Casolini F, Patti J, Patel P;
 PI Domanski P;
 XX WPI; 2001-607512/69.
 XX Novel isolated antibody which recognizes collagen-binding peptide such
 PT as CNA19 peptide from Staphylococcus aureus, useful for preventing or
 PT treating Staphylococcus aureus or epidermidis infection -
 XX Example 2; Fig 2A; 107pp; English.
 XX The invention relates to an antibody which recognises a collagen-binding
 CC region including CNA19 of CNA protein from Staphylococcus aureus. This
 CC antibody is cross-reactive to collagen binding region of both S. aureus
 CC and S. epidermidis. It is useful for preventing or treating S. aureus or
 CC S. epidermidis infection in human or animal, and for displacing S. aureus
 CC or S. epidermidis bound to collagen. Antibody of the invention is useful
 CC for interfering with, modulating, and inhibiting the binding interactions
 CC between Staphylococcal bacteria and collagen, for detecting the presence
 CC of Staphylococcal bacteria or Staphylococcal collagen or binding
 CC proteins, to diagnose Staphylococcal infection, as research tools, for
 CC development of vaccine for passive immunisation against Staphylococcal
 CC infections, and in production facilities or laboratories to isolate
 CC additional quantities of collagen-binding proteins. It is also useful
 CC for treating medical instruments in order to reduce or eliminate the
 CC possibility of them becoming infected or further spreading the
 CC infection. The present sequence is S. aureus CNA19 protein.
 XX
 XX Sequence 168 AA;

Query Match 90.5%; Score 19; DB 22; Length 168;
 Best Local Similarity 66.7%; Pred. No. 1e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
 ||| |
 Db 16 eagtss 21

RESULT 10
 AAR39711
 ID AAR39711 standard; Protein; 177 AA.

XX AC AAR39711;

XX DT 24-JAN-1994 (first entry)

XX DE A. oryzae WT neutral protease.

XX KW Wildtype; protease; variant; neutral; mercapto group; food products;
 XX KW thermal stability; soy fermentation.

XX OS Aspergillus oryzae.

XX PN JP05168479-A.

XX PD 02-JUL-1993.

XX PF 26-DEC-1991; 91JP-0344443.

XX CC

PR XX 26-DEC-1991; 91JP-0344443.
 XX (SHSA) SHOKUHN SANGYO KOSOKINO HENKA.
 XX WPI; 1993-247571/31.
 DR N-PSDB; AAQ46955.
 XX New variant neutral protease II - includes cysteine substd. with
 PT aminoacid having no mercapto gp. in aminoacid sequence of yellow
 PT green koji mould neutral protease II
 XX Disclosure; Page 7-8; 9pp; Japanese.
 XX This sequence represents a wildtype protease which may be used as the
 CC basis for the production of a variant neutral protease. The variant
 CC protease has either Cys6 or Cys78 substituted with an amino acid
 CC which has no -SH group. The variant proteases (see also AAR39713-14)
 CC have lower thermal stability than the WT and may be used in soy
 CC fermentation microorganisms. Soy produced by these microorganisms
 CC may be made into food products which will not be degraded by the
 CC presence of protease.
 XX
 XX Sequence 177 AA;

Query Match 90.5%; Score 19; DB 14; Length 177;
 Best Local Similarity 66.7%; Pred. No. 1.1e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
 ||| |
 Db 65 eagtss 70

RESULT 11
 AAR39712
 ID AAR39712 standard; Protein; 177 AA.

XX AC AAR39712;

XX DT 24-JAN-1994 (first entry)

XX DE A. oryzae C6A neutral protease.

XX KW Wildtype; protease; variant; neutral; mercapto group; food products;
 XX KW thermal stability; soy fermentation.

XX OS Aspergillus oryzae.

XX FH Key Location/Qualifiers
 XX FT Misc-difference 6
 XX FT /label= C6A

XX PN JP05168479-A.

XX PD 02-JUL-1993.

XX PF 26-DEC-1991; 91JP-0344443.

XX PR 26-DEC-1991; 91JP-0344443.

XX (SHSA) SHOKUHN SANGYO KOSOKINO HENKA.

XX WPI; 1993-247571/31.

XX New variant neutral protease II - includes cysteine substd. with
 PT aminoacid having no mercapto gp. in aminoacid sequence of yellow
 PT green koji mould neutral protease II
 XX Disclosure; Page 8; 9pp; Japanese.

XX The sequences given in AAR39712-13 represent variant neutral proteases
 CC based on the Aspergillus oryzae protease sequence (see also AAR39711).

CC These variant proteases have either Cys6 or Cys78 substituted with an
 CC amino acid which has no -SH group. These variant proteases have lower
 CC thermal stability than the WT and may be used in soy fermentation
 CC microorganisms. Soy produced by these microorganisms may be made into
 CC food products which will not be degraded by the presence of protease.

XX SQ Sequence 177 AA;

Query Match 90.5%; Score 19; DB 14; Length 177;
 Best Local Similarity 66.7%; Pred. No. 1.1e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 65 eagsts 70

RESULT 12

AAR39713
 ID AAR39713 standard; Protein; 177 AA.

XX AC AAR39713;

XX DT 24-JAN-1994 (first entry)

XX DE A. oryzae C78A neutral protease.

XX KW Wildtype; protease; variant; neutral; mercapto group; food products;
 KW thermal stability; soy fermentation.

XX OS Aspergillus oryzae.

XX FH Key Location/Qualifiers
 FT Misc-difference 78
 FT /label= C78A

XX PN JP05168479-A.

XX PD 02-JUL-1993.

XX PF 26-DEC-1991; 91JP-0344443.

XX PR 26-DEC-1991; 91JP-0344443.

XX PA (SHSA) SHOKUHIN SANGYO KOSOKINO HENKA.

XX DR WPI; 1993-247571/31.

XX PT New variant neutral protease II - includes cysteine substd. with
 PT aminoacid having no mercapto gp. in aminoacid sequence of yellow
 PT green koji mould neutral protease II

XX PS Disclosure; Page 8; 9pp; Japanese.

XX CC The sequences given in AAR39712-13 represent variant neutral proteases
 CC based on the Aspergillus oryzae protease sequence (see also AAR39711).
 CC These variant proteases have either Cys6 or Cys78 substituted with an
 CC amino acid which has no -SH group. These variant proteases have lower
 CC thermal stability than the WT and may be used in soy fermentation
 CC microorganisms. Soy produced by these microorganisms may be made into
 CC food products which will not be degraded by the presence of protease.

XX SQ Sequence 177 AA;

Query Match 90.5%; Score 19; DB 14; Length 177;
 Best Local Similarity 66.7%; Pred. No. 1.1e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 65 eagsts 70

RESULT 13

AAB24437
 ID AAB24437 standard; Protein; 207 AA.

XX AC AAB24437;

XX DT 20-NOV-2000 (first entry)

XX DE Human secreted protein sequence encoded by gene 1 SEQ ID NO:62.

XX KW Human; secreted protein; cytostatic; antianaemic; antidiabetic;
 KW antiinflammatory; ophthalmological; antirheumatic; antiarthritic;
 KW antipsoriatic; antiangiogenic; cardiant; anti-HIV; nootropic;
 KW neuroprotective; antimicrobial; antiparkinsonian; cancer;
 KW immune system disorder; angiogenesis; hyperproliferative disorder;
 KW cardiovascular disorder; apoptosis; neuroproliferative disease;
 KW infectious disease; wound healing.

XX OS Homo sapiens.

XX PN WO200035937-A1.

XX PD 22-JUN-2000.

XX PF 16-DEC-1999; 99WO-US29950.

XX PR 17-DEC-1998; 98US-0112809.

XX PR 18-DEC-1998; 98US-0113006.

XX PA (HUMA-) HUMAN GENOME SCI INC.

XX PI Ruben SM, Ebner R, Rosen CA, Endress GA, Soppet DR, Ni J;

PI Duan DR, Moore PA, Shi Y, Lafleur DW, Olsen HS, Florence K;

XX DR WPI; 2000-431566/37.

XX DR N-PSDB; AAA78381.

XX PT Forty seven human nucleic acids encoding secreted proteins, useful in
 PT the treatment, prevention and diagnosis of cancers, disorders of the
 PT immune system, angiogenesis disorders, neurological diseases and
 PT hyperproliferative disorders -

XX PS Claim 11; Page 478-479; 562pp; English.

XX CC The polynucleotide sequence given in AAA78381 to AAA78432 encode the
 CC human secreted proteins given in AAB24437 to AAB24604. Human secreted
 CC proteins have activities based on the tissues and cells the genes are
 CC expressed in. Examples of activities include: cytostatic; antianaemic;
 CC antidiabetic; antiinflammatory; ophthalmological; antirheumatic;
 CC antiarthritic; antipsoriatic; antiangiogenic; cardiant; anti-HIV;
 CC nootropic; neuroprotective; antimicrobial and antiparkinsonian.
 CC Human secreted protein polynucleotides, polypeptides, antagonists and/or
 CC agonists may be useful in treating, preventing, and/or diagnosing other
 CC diseases, disorders, and/or conditions such as: (a) cancers; (b)
 CC disorders of the immune system; (c) angiogenesis disorders; (d)
 CC hyperproliferative disorders; (e) cardiovascular disorders; (f) diseases
 CC associated with increase apoptosis; (g) neurological diseases; and
 CC (h) infectious diseases. They are also used to promote wound healing.
 CC AAA78372 to AAA78380 and AAB24436 represent sequences used in the
 CC exemplification of the present invention.

XX SQ Sequence 207 AA;

Query Match 90.5%; Score 19; DB 21; Length 207;
 Best Local Similarity 66.7%; Pred. No. 1.3e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 77 eagats 82

RESULT 14
 ID AAW31553 standard; Protein; 211 AA.
 XX AC AAW31553;
 XX DT 21-MAY-1998 (first entry)
 XX DE Collagen binding protein M31 epitope.
 XX KW Collagen binding protein; cna gene; sepsis; infection;
 KW microbial surface component recognising adhesive matrix molecule;
 KW MSCRAMM; adhesin; vaccine; immunisation; diagnosis; therapy;
 KW epitope M31.
 XX OS Staphylococcus aureus.
 XX FH Key Location/Qualifiers
 FT Peptide 1..12
 FT Protein /note= "vector pQE30-derived peptide"
 FT 13..211
 FT /note= "epitope M31"
 XX WO9743314-A2.
 XX PN 20-NOV-1997.
 XX PD 14-MAY-1997; 97WO-US08210.
 XX PF 16-MAY-1996; 96US-0017678.
 XX PR (UABR-) UAB RES FOUND.
 XX PA (TEXA) UNIV TEXAS A & M SYSTEM.
 XX PI Hook M, House-Pompeo K, Patti JM, Sthanam N, Symersky J;
 XX WPI; 1998-008801/01.
 XX DR N-PSDB; AAT93437.
 XX PT Antibody that interacts with collagen binding domain of
 PT Staphylococcal cna gene product - useful to prevent bacterial sepsis
 PT in animal infected with Staphylococcus aureus
 XX Claim 31; Page 115-116; 143pp; English.
 XX This protein comprises Staphylococcus aureus collagen binding
 CC protein (CBP) epitope M31, i.e. amino acids 61-343 of full-length
 CC CBP, plus a vector-derived N-terminal peptide. Claimed 441, 849
 CC and 1500 bp nucleic acid sequences (see AAT93436-38) respectively
 CC encode CBP epitopes M17, M31 and M55 (see AAW31552-54) that confer
 CC protection against S. aureus infection. These nucleic acid
 CC sequences can be used in the recombinant production of the CBP
 CC epitopes. The CBP protein and antigenic epitopes are contemplated
 CC for use in the treatment of pathological infections, especially to
 CC prevent bacterial adhesion to collagen. The claimed nucleic acids
 CC as well as claimed anti-CBP antibodies will also be of use in
 CC screening, diagnostic and therapeutic applications including active
 CC and passive immunisation and methods for the prevention of
 CC bacterial colonisation in an animal such as a human. The CBP
 CC epitopes are also contemplated for use in the preparation of
 CC vaccines and as carrier proteins in vaccine formulations, as well
 CC as in the formulation of compositions for the prevention of S.
 CC aureus infection.
 XX Sequence 211 AA;
 XX SQ
 Query Match 90.5%; Score 19; DB 19; Length 211;
 Best Local Similarity 66.7%; Pred. No. 1.3e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 Db 34 eagtss 39
 RESULT 15
 ID AAW31553 standard; Protein; 225 AA.
 XX AC AAW31553;
 XX DT 13-FEB-2002 (first entry)
 XX DE Novel human diagnostic protein #3042.
 XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder.
 XX OS Homo sapiens.
 XX FH Key Location/Qualifiers
 FT Peptide 1..12
 FT Protein /note= "vector pQE30-derived peptide"
 FT 13..211
 FT /note= "epitope M31"
 XX WO9743314-A2.
 XX PN 20-NOV-1997.
 XX PD 14-MAY-1997; 97WO-US08210.
 XX PF 16-MAY-1996; 96US-0017678.
 XX PR (UABR-) UAB RES FOUND.
 XX PA (TEXA) UNIV TEXAS A & M SYSTEM.
 XX PI Hook M, House-Pompeo K, Patti JM, Sthanam N, Symersky J;
 XX WPI; 1998-008801/01.
 XX DR N-PSDB; AAT93437.
 XX PT Antibody that interacts with collagen binding domain of
 PT Staphylococcal cna gene product - useful to prevent bacterial sepsis
 PT in animal infected with Staphylococcus aureus
 XX Claim 31; Page 115-116; 143pp; English.
 XX This protein comprises Staphylococcus aureus collagen binding
 CC protein (CBP) epitope M31, i.e. amino acids 61-343 of full-length
 CC CBP, plus a vector-derived N-terminal peptide. Claimed 441, 849
 CC and 1500 bp nucleic acid sequences (see AAT93436-38) respectively
 CC encode CBP epitopes M17, M31 and M55 (see AAW31552-54) that confer
 CC protection against S. aureus infection. These nucleic acid
 CC sequences can be used in the recombinant production of the CBP
 CC epitopes. The CBP protein and antigenic epitopes are contemplated
 CC for use in the treatment of pathological infections, especially to
 CC prevent bacterial adhesion to collagen. The claimed nucleic acids
 CC as well as claimed anti-CBP antibodies will also be of use in
 CC screening, diagnostic and therapeutic applications including active
 CC and passive immunisation and methods for the prevention of
 CC bacterial colonisation in an animal such as a human. The CBP
 CC epitopes are also contemplated for use in the preparation of
 CC vaccines and as carrier proteins in vaccine formulations, as well
 CC as in the formulation of compositions for the prevention of S.
 CC aureus infection.
 XX Sequence 225 AA;
 XX SQ
 Query Match 90.5%; Score 19; DB 22; Length 225;
 Best Local Similarity 66.7%; Pred. No. 1.4e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```
Db          ||| . |
            16 eagtss 21

RESULT 16
AAG05085
ID  AAG05085 standard; Protein; 229 AA.
XX
AC  AAG05085;
XX
DT  17-OCT-2000 (first entry)
XX
DE  Arabidopsis thaliana protein fragment SEQ ID NO: 1356.
XX
KW  Protein identification; signal transduction pathway; metabolic pathway;
KW  hybridisation assay; genetic mapping; gene expression control; promoter;
KW  termination sequence.
XX
OS  Arabidopsis thaliana.
XX
EP  EP1033405-A2.
XX
PD  06-SEP-2000.
XX
PF  25-FEB-2000; 2000EP-0301439.
XX
PR  25-FEB-1999; 99US-0121825.
PR  05-MAR-1999; 99US-0123180.
PR  09-MAR-1999; 99US-0123548.
PR  23-MAR-1999; 99US-0125788.
PR  25-MAR-1999; 99US-0126264.
PR  29-MAR-1999; 99US-0126785.
PR  01-APR-1999; 99US-0127462.
PR  06-APR-1999; 99US-0128234.
PR  08-APR-1999; 99US-0128714.
PR  16-APR-1999; 99US-0129845.
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Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 203 eagsss 208

RESULT 18
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AC AAG05084;
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DT 17-OCT-2000 (first entry)
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KW Protein identification; signal transduction pathway; metabolic pathway;
KW hybridisation assay; genetic mapping; gene expression control; promoter;
KW termination sequence.
XX
OS Arabidopsis thaliana.
XX
PN EPI033405-A2.
XX
PD 06-SEP-2000.
XX
PF 25-FEB-2000; 2000EP-0301439.
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Best Local Similarity 66.7%; Pred. No. 1.5e+03;

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Db 170 eaqsss 175

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AC AA95431;
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DT 10-OCT-2000 (first entry)
XX
DE Human calcium channel SOC-2/CRAC-1.
XX
KW SOC-2/CRAC-1; calcium channel; human; store operated channel;
KW calcium release activated channel; therapy; diagnosis;
KW lymphocyte proliferative disorder.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Misc-difference 103 /note= "encoded by SCA"
FT Misc-difference 104 /note= "encoded by CSA"
FT Misc-difference 105 /note= "encoded by RSC"
FT Misc-difference 109 /note= "encoded by GNT"
FT Misc-difference 141 /note= "encoded by NCA"
PN WO200040614-A2.
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PD 13-JUL-2000.
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PF 20-DEC-1999; 99WO-US29996.
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PR 22-JUN-1999; 99US-0140415.
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PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
XX
PI Scharenberg AM;
XX
XX WPI; 2000-465957/40.
DR N-PSDB; AAA49918.
XX
PT New SOC/CRAC calcium channel polynucleotides and polypeptides used to
PT diagnose and treat proliferative disorders associated with the channel,
PT and to screen for novel modulators of the channel -
XX
PS Claim 14; Page 58-59; 108pp; English.
XX
CC The present sequence is that of a partial sequence of human
CC SOC-2/CRAC-1 (full-length sequence given in AA95431), as deduced
CC from a partial cDNA clone (see AAA49918). SOC-2/CRAC-1 is a member
CC of a novel family of store operated channel (SOC) or calcium release
CC activated channel (CRAC) polypeptides that modulate Ca²⁺ flux into
CC and out of a cell, and which may be activated upon depletion of
CC Ca²⁺ from intracellular calcium stores, allowing Ca²⁺ influx into
CC a cell. SOC-2/CRAC-1 is expressed predominantly in human
CC haematopoietic cells, liver, spleen, heart and kidney.
CC Compositions for expressing functional SOC/CRAC calcium channel
CC polypeptides in cells are expected to be useful for treating
CC patients that have reduced extracellular calcium influx into their
CC SOC/CRAC-expressing cells. They will also be useful for delivering
CC therapeutic and/or imaging agents to such cells to modulate
CC proliferation and growth. SOC/CRAC polypeptides also represent
CC targets for designing and/or identifying inhibitors that block
CC lymphocyte proliferation and binding agents that selectively bind
CC to SOC/CRAC polypeptides to which drugs or toxins can be conjugated
CC for delivery to SOC/CRAC expression in a subject can be used to assess
CC the level of SOC/CRAC expression in a subject can be used to assess
CC the presence, or absence, or stage of a proliferative disorder,

CC e.g. a lymphocyte proliferative disorder.

XX Sequence 243 AA;

Query Match 90.58; Score 19; DB 21; Length 243;
Best Local Similarity 66.78; Pred. No. 1.5e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

Db 53 eagsss 58

RESULT 20

AAG05083

ID AAG05083 standard; Protein; 257 AA.

AC AAG05083;

XX 17-OCT-2000 (first entry)

XX Arabidopsis thaliana protein fragment SEQ ID NO: 1354.

XX Protein identification; signal transduction pathway; metabolic pathway;
KW hybridisation assay; genetic mapping; gene expression control; promoter;
KW termination sequence.

XX Arabidopsis thaliana.

PN EP1033405-A2.

XX 06-SEP-2000.

XX 25-FEB-2000; 2000EP-0301439.

XX 25-FEB-1999; 99US-0121825.

XX 03-MAR-1999; 99US-0123180.

XX 09-MAR-1999; 99US-0123548.

XX 23-MAR-1999; 99US-0125788.

XX 25-MAR-1999; 99US-0126264.

XX 29-MAR-1999; 99US-0126785.

XX 01-APR-1999; 99US-0127462.

XX 08-APR-1999; 99US-0128234.

XX 16-APR-1999; 99US-0128714.

XX 19-APR-1999; 99US-0129845.

XX 21-APR-1999; 99US-0130077.

XX 23-APR-1999; 99US-0130510.

XX 23-APR-1999; 99US-0130891.

XX 28-APR-1999; 99US-0131149.

XX 30-APR-1999; 99US-0132048.

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XX 05-MAY-1999; 99US-0132484.

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PR 28-JUL-1999; 99US-0145951.
PR 02-AUG-1999; 99US-0146386.
PR 02-AUG-1999; 99US-0146388.
PR 02-AUG-1999; 99US-0146389.
PR 03-AUG-1999; 99US-0147038.
PR 04-AUG-1999; 99US-0147204.
PR 04-AUG-1999; 99US-0147302.
PR 05-AUG-1999; 99US-0147192.
PR 05-AUG-1999; 99US-0147260.

		Matches	4;	Conservative	0;	Mismatches	2;	Indels	0;	Gaps	0;
Qy	1	eaqxss	6								
Db	184	eaqsss	189								
RESULT 21											
AAU68590											
ID	AAU68590	standard; Protein; 264 AA.									
XX	AAU68590;										
XX	16-JAN-2002	(first entry)									
DE	Human novel cytokine encoded by cDNA 790CIP2D_8 #2.										
KW	Human; cytokine; cell proliferation; cell differentiation; antiinflammatory; stem cell growth factor; activin; inhibin; cancer; nervous system disease; neuropathy; Alzheimer's disease; Parkinson's disease; Huntington's disease; spinal cord disorder; head trauma; stroke; myeloid cell disorder; lymphoid cell disorder; platelet disorder; thrombocytopaenia; stem cell disorder; aplastic anaemia; tissue regeneration; wound healing; ulcer; osteoporosis; osteoarthritis; bone degenerative disorder; periodontal disease; fibrosis; reperfusion; immune disorder; SCID; severe combined immunodeficiency; infection; autoimmune disorder; multiple sclerosis; rheumatoid arthritis; diabetes mellitus; allergy; asthma; coagulation disorder; haemophilia; sepsis; nephritis; inflammatory bowel disease; food supplement; immunogen.										
XX	Homo sapiens.										
XX	WO200175093-A1.										
XX	11-OCT-2001.										
XX	30-MAR-2001; 2001WO-US10484.										
XX	31-MAR-2000; 2000US-0540217.										
PR	23-AUG-2000; 2000US-0649167.										
PR	22-SEP-2000; 2000US-0668680.										
PR	23-OCT-2000; 2000US-0695618.										
PR	30-NOV-2000; 2000US-0728711.										
PR	14-MAR-2001; 2000US-0728711.										
XX	(HYSE-) HYSEQ INC.										
XX	Tang YT, Asundi V, Zhou P, Xue AJ, Ren F, Zhang J, Wang J, Xu C; Yang Y, Zabo QA, Chen R, Wang D, Goodrich RW, Liu C, Drmanac RT; WPI; 2001-626432/72.										
DR	N-PSDB; AAS59882.										
XX	New polypeptides and nucleic acids, useful for diagnosis, treatment of inflammatory, autoimmune, neurological, myeloid or lymphoid cell, bone degenerative disorders, cancer and promoting wound healing										
PS	Claim 20; Page 328; 336pp; English.										
XX	The invention relates to isolated human polypeptides (which may be cytokines) and the polynucleotides encoding them. The protein is useful for identifying a compound which binds to it (e.g. modulators, agonists and antagonists). The polynucleotides are useful as an array for mismatch detection. The proteins and nucleic acids are useful as nutritional sources or supplements. The protein exhibits activity relating to cytokine, cell proliferation, cell differentiation, antiinflammatory, stem cell growth factor activity, immune stimulating or immune suppressing and activin or inhibin related activities. The proteins (and antibodies raised against them) and nucleic acids are therefore useful in the diagnosis and treatment of diseases and disorders such as cancer, central and peripheral nervous system diseases and neuropathies, Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic										

CC lateral sclerosis, spinal cord disorders, head trauma, cerebrovascular
 CC diseases, stroke, myeloid or lymphoid cell disorders, platelet disorders,
 CC thrombocytopenia, stem cell disorders, aplastic anaemia, for
 CC regeneration of bone, cartilage, tendon, ligament and/or nerve tissue
 CC growth, and in tissue repair, healing of burns, incisions, ulcers, for
 CC treating osteoporosis, osteoarthritis, bone degenerative disorders, or
 CC periodontal disease, lung or liver fibrosis, reperfusion injury in
 CC various tissues, various immune deficiencies and disorders including
 CC severe combined immunodeficiency (SCID), bacterial or fungal infections,
 CC autoimmune disorders (e.g. multiple sclerosis, rheumatoid arthritis,
 CC diabetes mellitus, myasthenia gravis), allergic reactions and conditions,
 CC such as asthma or other respiratory problems, coagulation disorders,
 CC haemophilia), septic shock, sepsis, arthritis, nephritis and inflammatory
 CC bowel disease, viral infection and are useful in altering bodily
 CC characteristics. The present sequence represents a novel protein of the
 CC invention.

XX Sequence 264 AA;

Query Match 90.5%; Score 19; DB 22; Length 264;

Best Local Similarity 66.7%; Pred. No. 1.6e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6

||| |

Db 207 eagaas 212

RESULT 22

ABB69721
 ID ABB69721 standard; Protein; 270 AA.

XX AC ABB69721;

XX DT 26-MAR-2002 (first entry)

XX DE Drosophila melanogaster polypeptide SEQ ID NO 35955.

XX KW Drosophila; developmental biology; cell signalling; insecticide;

XX KW pharmaceutical.

XX OS Drosophila melanogaster.

XX PN WO200171042-A2.

XX PD 27-SEP-2001.

XX PF 23-MAR-2001; 2001WO-US09231.

XX PR 23-MAR-2000; 2000US-191637P.

XX PR 11-JUL-2000; 2000US-0614150.

XX PA (PEKE) PE CORP NY.

XX PI Venter JC, Adams M, Li PWD, Myers EW;

XX DR WPI; 2001-656860/75.

XX DR N-PSDB; ABL13824.

XX PT New isolated nucleic acid detection reagent for detecting 1000 or more
 PT genes from Drosophila and for elucidating cell signalling and cell-cell
 PT interactions -

XX PS Disclosure; SEQ ID NO 35955; 21pp + Sequence Listing; English.

XX CC The invention relates to an isolated nucleic acid detection reagent
 CC capable of detecting 1000 or more genes from Drosophila. The invention is
 CC useful in developmental biology and in elucidating cell signalling and
 CC cell-cell interactions in higher eukaryotes for the development of
 CC insecticides, therapeutics and pharmaceutical drugs. The invention
 CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
 CC sequences (ABL01840-ABL16175) and the encoded proteins

CC (ABB57737-ABB72072).

CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 270 AA;

Query Match 90.5%; Score 19; DB 22; Length 270;

Best Local Similarity 66.7%; Pred. No. 1.7e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6

||| |

Db 169 eagsts 174

RESULT 23

ABG03786

ID ABG03786 standard; Protein; 270 AA.

XX AC ABG03786;

XX DT 13-FEB-2002 (first entry)

XX DE Novel human diagnostic protein #3777.

XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

XX KW food supplement; medical imaging; diagnostic; genetic disorder.

XX OS Homo sapiens.

XX PN WO200175067-A2.

XX PD 11-OCT-2001.

XX PF 30-MAR-2001; 2001WO-US08631.

XX PR 31-MAR-2000; 2000US-0540217.

XX PR 23-AUG-2000; 2000US-0649167.

XX PA (HYSE-) HYSEQ INC.

XX PI Drmanac RT, Liu C, Tang YT;

XX DR WPI; 2001-639362/73.

XX DR N-PSDB; AAS67973.

XX PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -

XX PS Claim 20; SEQ ID NO 34145; 103pp; English.

XX CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human
 CC diagnostic amino acid sequences of the invention.

CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

SQ Sequence 270 AA;

Query Match 90.5%; Score 19; DB 22; Length 270;

Best Local Similarity 66.7%; Pred. No. 1.7e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

Db 16 eagtss 21

RESULT 24

AAU311144

ID AAU311144 standard; Protein; 273 AA.

XX AC AAU311144;

XX DT 18-DEC-2001 (first entry)

XX DE Novel human secreted protein #1635.

XX KW Human; vaccination; gene therapy; nutritional supplement;

XX KW stem cell proliferation; haematopoiesis; nerve tissue regeneration;

XX KW immune suppression; immune stimulation; anti-inflammatory; leukaemia.

XX OS Homo sapiens.

XX PN WO200179449-A2.

XX PD 25-OCT-2001.

XX PF 16-APR-2001; 2001WO-US08656.

XX PR 18-APR-2000; 2000US-0552929.

XX PR 26-JAN-2001; 2001US-0770160.

XX PA (HYSE-) HYSEQ INC.

XX PI Tang YT, Liu C, Drmanac RT;

XX WPI; 2001-611725/70.

XX PT Nucleic acids encoding a range of human polypeptides, useful in genetic

XX PT vaccination, testing and therapy -

XX PS Claim 20; Page 409; 765pp; English.

XX CC The invention relates to novel human secreted polypeptides. The

XX CC polypeptides and antibodies to the polypeptides are useful for

XX CC determining the presence of or predisposition to a disease associated

XX CC with altered levels of polypeptide. The polypeptides are also useful for

XX CC identifying agents (agonists and antagonists) that bind to them. Cells

XX CC expressing the proteins are useful for identifying a therapeutic agent

XX CC for use in treatment of a pathology related to aberrant expression or

XX CC physiological interactions of the polypeptide. Vectors comprising

XX CC the nucleic acids encoding the polypeptides and cells genetically

XX CC engineered to express them are also useful for producing the proteins.

XX CC The proteins are useful in genetic vaccination, testing and

XX CC therapy, and can be used as nutritional supplements. They may be used to

XX CC increase stem cell proliferation; to regulate haematopoiesis; and in

XX CC bone, cartilage, tendon and/or nerve tissue growth or regeneration;

XX CC immune suppression and/or stimulation; as anti-inflammatory agents; and

XX CC in treatment of leukaemias. AAU29510-AAU33304 represent the amino acid

XX CC sequences of novel human secreted proteins of the invention.

XX SQ Sequence 273 AA;

Query Match

Best Local Similarity 90.5%; Score 19; DB 22; Length 273;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

Db 35 eagass 40

RESULT 25

AAU47789

ID AAU47789 standard; Protein; 290 AA.

XX AC AAU47789;

XX DT 27-FEB-2002 (first entry)

XX DE Propionibacterium acnes immunogenic protein #8685.

XX KW SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;

XX KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;

XX KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;

XX KW dermatological; osteopathic; neuroprotectant.

XX OS Propionibacterium acnes.

XX PN WO200181581-A2.

XX PD 01-NOV-2001.

XX PF 20-APR-2001; 2001WO-US12865.

XX PR 21-APR-2000; 2000US-199047P.

XX PR 02-JUN-2000; 2000US-208841P.

XX PR 07-JUL-2000; 2000US-216747P.

XX PA (CORI-) CORIXA CORP.

XX PI Skeiky YAW, Persing DH, Mitcham JL, Wang SS, Bhatia A;

XX PI L'maisonneuve J, Zhang Y, Jen S, Carter D;

XX WPI; 2001-616774/71.

XX N-PSDB; AAS59539.

XX PT Propionibacterium acnes polypeptides and nucleic acids useful for

XX PT vaccinating against and diagnosing infections, especially useful for

XX PT treating acne vulgaris -

XX PS Example 1; SEQ ID No 8984; 1069pp; English.

XX CC Sequences AAU39105-AAU68017 represent Propionibacterium acnes immunogenic

XX CC polypeptides. The proteins and their associated DNA sequences are used in

XX CC the treatment, prevention and diagnosis of medical conditions caused by

XX CC P. acnes. The disorders include SAPHO syndrome (synovitis, acne,

XX CC pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.

XX CC P. acnes is also involved in infections of bone, joints and the central

XX CC nervous system, however it is particularly involved in the inflammatory

XX CC lesions associated with acne vulgaris. A method for detecting the

XX CC presence or absence of P. acnes in a patient comprises contacting a

XX CC sample with a binding agent that binds to the proteins of the invention

XX CC and determining the amount of bound protein in the sample. The

XX CC polypeptides may be used as antigens in the production of antibodies

XX CC specific for P. acnes proteins. These antibodies can be used to

XX CC downregulate expression and activity of P. acnes polypeptides and

XX CC therefore treat P. acnes infections. The antibodies may also be used as

XX CC diagnostic agents for determining P. acnes presence, for example, by

XX CC enzyme linked immunosorbent assay (ELISA).

XX CC Note: The sequence data for this patent did not form part of the printed

XX CC specification, but was obtained in electronic format directly from WIPO

XX CC at ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 290 AA;

Query Match 90.5%; Score 19; DB 22; Length 290;
Best Local Similarity 66.7%; Pred. No. 1.8e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
| | | |
Db 79 eagass 84

RESULT 26

ABG02081
ID ABG02081 standard; Protein; 305 AA.

XX AC ABG02081;

XX DT 13-FEB-2002 (first entry)

XX DE Novel human diagnostic protein #2072.

XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.

XX OS Homo sapiens.

XX PN WO200175067-A2.

XX PD 11-OCT-2001.

XX PF 30-MAR-2001; 2001WO-US08631.

XX PR 31-MAR-2000; 2000US-0340217.

XX PR 23-AUG-2000; 2000US-0649167.

XX PA (HYSE-) HYSEQ INC.

XX PI Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

XX N-PSDB; AAS66268.

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -

PS Claim 20; SEQ ID No 32440; 103pp; English.

CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.

CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 305 AA;

SQ

Query Match 90.5%; Score 19; DB 22; Length 305;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
| | | |
Db 178 eagass 183

RESULT 27

AAB94964

ID AAB94964 standard; Protein; 323 AA.

XX AC AAB94964;

XX DT 26-JUN-2001 (first entry)

XX DE Human protein sequence SEQ ID NO:16523.

XX KW Human; primer; detection; diagnosis; antisense therapy; gene therapy.

XX OS Homo sapiens.

XX PN EP1074617-A2.

XX PD 07-FEB-2001.

XX PF 28-JUL-2000; 2000EP-0116126.

XX PR 29-JUL-1999; 99JP-0248036.

XX PR 27-AUG-1999; 99JP-0300253.

XX PR 11-JAN-2000; 2000JP-0118776.

XX PR 02-MAY-2000; 2000JP-0183767.

XX PR 09-JUN-2000; 2000JP-0241899.

XX PA (HELI-) HELIX RES INST.

XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

XX PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX WPI; 2001-318749/34.

PT Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs -

PS Claim 8; SEQ ID 16523; 2537pp + CD ROM; English.

CC The present invention describes primer sets for synthesising 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesising polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.

SQ Sequence 323 AA;

Query Match 90.5%; Score 19; DB 22; Length 323;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 308 eagsts 313

RESULT 28
AAR05528
ID AAR05528 standard; protein; 330 AA.
XX
AC AAR05528;
XX
DT 23-OCT-1990 (first entry)
XX
DE High density lipoprotein (HDL) binding protein.
XX
KW High density lipoprotein; HDL-binding protein; atherosclerosis;
KW hypercholesterolaemia; ds.
XX
OS Homo sapiens.
XX
PN WO9005744-A.
XX
PO 31-MAY-1990.
XX
PF 17-NOV-1989; 89WO-0005169.
XX
PR 18-NOV-1988; 88US-0273388.
XX
PA (UNIW) UNIV OF WASHINGTON.
XX
PA (ZYMO-) ZYMOGENETICS INC.
XX
PI Oram JF, McKnight GL, Hart CE, Curtis DA;
XX
DR WPI; 1990-193405/25.
DR N-PSDB; AAQ04784.
XX
PT New mammalian proteins binding high density lipoprotein sub-class 3 -
PT DNA encoding them and derived antibodies, for screening
PT potentially therapeutic HDL analogues and for diagnosing risk of
PT atherosclerosis.
XX
PS Claim 4; Fig 1A-D; 79pp; English.
XX
CC The protein product may be used to raise Abs, and the cDNA to
CC create probes, both useful in screening for HDL analogues,
CC agonists and antagonists, and in identifying abnormalities in the
CC HDL binding/receptor pathway. HDL analogues can be used in treating
CC hypercholesterolaemia and atherosclerosis
XX
SQ Sequence 330 AA;

Query Match 90.5%; Score 19; DB 11; Length 330;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 205 eagas 210

RESULT 29
AAY95898
ID AAY95898 standard; Protein; 332 AA.
XX
AC AAY95898;
XX

20-NOV-2000 (first entry)
Human myristoylated alanine-rich C kinase substrate MARCKS.
XX
DE MARCKS; myristoylated alanine-rich C kinase substrate; human;
XX mucus secretion; inhibitor; bronchitis; asthma; cystic fibrosis;
KW chronic obstructive pulmonary disease; pneumonia; emphysema;
KW influenza; rhinitis; therapy.
XX
OS Homo sapiens.
XX
XX
FH Key Location/Qualifiers
FT Misc-difference 84 /note= "Ser in sequence of AAY95899"
FT Misc-difference 119 /note= "Ala in sequence of AAY95899"
FT Peptide 2...25
FT /note= "MANS peptide of AAY95896"
FT Peptide 152...176
FT /note= "NA-PSD peptide of AAY95897"
XX
PN WO200050062-A2.
XX
XX
PD 31-AUG-2000.
XX
XX
PF 24-FEB-2000; 2000WO-US05050.
XX
PR 24-FEB-1999; 99US-0256154.
XX
PA (UYNC-) UNIV NORTH CAROLINA STATE.
XX
PI Li Y, Martin LD, Adler KB;
XX
DR WPI: 2000-572036/53.
DR N-PSDB; AAR50339.
XX
PT Regulating mucus secretion by a mucus-secreting cell, useful for
PT treating e.g. bronchitis, asthma or pneumonia, by administering a
PT compound that inhibits or enhances myristolated alanine-rich C-kinase
PT substrate protein -
XX
PS Claim 3; Page 42-43; 66pp; English.
XX
CC The present sequence is that of human myristoylated alanine-rich C
CC kinase substrate MARCKS protein, a major cellular substrate. The
CC invention relates to methods of inhibiting mucus secretion by a
CC mucus-secreting cell by administering a compound that inhibits
CC MARCKS protein-related mucus secretion. Such compounds include
CC active fragments of MARCKS protein such as MANS peptide (see
CC AAY95897) and NA-PSD peptide (see AAY95897), which corresponds to a
CC phosphorylation site of MARCKS. The inhibitor compounds can be
CC used to treat conditions such as bronchitis, cystic fibrosis,
CC chronic obstructive pulmonary disease, asthma, emphysema,
CC pneumonia, influenza, rhinitis and the common cold. An alternative
CC sequence for MARCKS is provided in AAY95899, which differs from the
CC present sequence at 2 amino acid residues, Ala-84 (Ser) and
CC Pro-119 (Ala).
XX
SQ Sequence 332 AA;

Query Match 90.5%; Score 19; DB 21; Length 332;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 206 eagas 211

RESULT 30
AAY95899
ID AAY95899 standard; Protein; 332 AA.

XX 26-SEP-2001 (first entry)
 XX C glutamicum protein fragment SEQ ID NO: 4543.
 DE Coryneform bacterium; amino acid synthesis; vitamin; saccharide;
 KW organic acid synthesis.
 KW Corynebacterium glutamicum.
 OS EP1108790-A2.
 PN 20-JUN-2001.
 XX 18-DEC-2000; 2000EP-0127688.
 PF 16-DEC-1999; 99JP-0377484.
 PR 07-APR-2000; 2000JP-0159162.
 PR 03-AUG-2000; 2000JP-0280988.
 XX (KYOWA) KYOWA HAKKO KOGYO KK.
 PA Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;
 PI Tateishi N, Senoh A, Ikeda M, Ozaki A;
 PI WPI; 2001-376931/40.
 DR N-PSDB; AAH66008.
 XX Novel polynucleotides derived from Coryneform bacteria, for identifying
 PT mutation point of a gene, measuring expression of a gene, analysing
 PT expression profile or pattern of a gene and identifying homologous gene
 PT -
 PS Claim 17; SEQ ID NO: 4543; 246pp + Sequence Listing; English.
 XX The present invention provides a number of nucleotide and protein
 CC sequences from the Coryneform bacterium Corynebacterium glutamicum. These
 CC are useful for identifying the mutation point of a gene derived from a
 CC mutant of coryneform bacterium, measuring expression amount and
 CC analysing the expression profile or expression pattern of a gene derived
 CC from Coryneform bacterium, and identifying a homologue of a gene derived
 CC from coryneform bacterium. Coryneform bacteria are useful for producing
 CC amino acids, nucleic acids, vitamins, saccharides and organic acids,
 CC particularly L-lysine. The present sequence is a protein described
 CC in the exemplification of the invention.
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from the
 CC European Patent Office.
 XX Sequence 341 AA;
 SQ

Query Match 90.5%; Score 19; DB 22; Length 341;
 Best Local Similarity 66.7%; Pred. No. 2.le+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 eagxxs 6
 ||| |
 Db 239 eagtss 244

RESULT 33
 AAB79110
 ID AAB79110 standard; Protein; 341 AA.
 XX AAB79110;
 AC AAB79110;
 XX 30-APR-2001 (first entry)
 DT Corynebacterium glutamicum HA protein sequence SEQ ID NO:176.
 DE Corynebacterium glutamicum; homeostasis; adaptation; HA protein;
 XX fine chemical production; organic acid; proteinogenic amino acid;
 KW

KW nonproteinogenic amino acid; purine base; pyrimidine base; nucleoside;
 KW nucleotide; lipid; saturated fatty acid; unsaturated fatty acid; diol;
 KW carbohydrate; aromatic compound; vitamin; cofactor; polyketide; enzyme;
 KW diagnosis; Corynebacterium diphtheriae; genetic engineering;
 KW Brevibacterium; environmental condition.
 XX Corynebacterium glutamicum.
 OS WO200100842-A2.
 PN 04-JAN-2001.
 XX 23-JUN-2000; 2000WO-IB00911.
 PF 25-JUN-1999; 99US-0141031.
 XX 08-JUL-1999; 99DE-1031636.
 PR 09-JUL-1999; 99DE-1032125.
 PR 09-JUL-1999; 99DE-1032126.
 PR 09-JUL-1999; 99DE-1032127.
 PR 09-JUL-1999; 99DE-1032128.
 PR 09-JUL-1999; 99DE-1032129.
 PR 09-JUL-1999; 99DE-1032226.
 PR 14-JUL-1999; 99DE-1032920.
 PR 14-JUL-1999; 99DE-1032922.
 PR 14-JUL-1999; 99DE-1032924.
 PR 14-JUL-1999; 99DE-1032928.
 PR 14-JUL-1999; 99DE-1032930.
 PR 14-JUL-1999; 99DE-1032933.
 PR 14-JUL-1999; 99DE-1032935.
 PR 14-JUL-1999; 99DE-1032973.
 PR 14-JUL-1999; 99DE-1033002.
 PR 14-JUL-1999; 99DE-1033003.
 PR 14-JUL-1999; 99DE-1033005.
 PR 14-JUL-1999; 99DE-1033006.
 PR 31-AUG-1999; 99DE-1041378.
 PR 31-AUG-1999; 99DE-1041379.
 PR 31-AUG-1999; 99DE-1041390.
 PR 31-AUG-1999; 99DE-1041391.
 PR 03-SEP-1999; 99DE-1042088.
 XX (BADI) BASF AG.
 XX Pompejus M, Kroeger B, Schroeder H, Zelder O, Haberhauer G;
 WPI; 2001-061974/07.
 DR N-PSDB; AAF71225.
 XX New isolated Corynebacterium glutamicum nucleic acid for production or
 PT modulation of production of fine chemicals such as amino acids,
 PT nucleosides, nucleotides, lipids, fatty acids, carbohydrates, vitamins
 PT or enzymes -
 XX Claim 20; Page 383-384; 712pp; English.
 PS AAF71138 to AAF71357 encode the Corynebacterium glutamicum homeostasis
 CC and adaptation (HA) proteins given in AAB79023 to AAB79242. The
 CC C. glutamicum HA genes (I) can be used in vectors for expression in host
 CC cells and production of fine chemicals, such as, an organic acid,
 CC proteinogenic or nonproteinogenic amino acid (preferred), purine or
 CC pyrimidine base, nucleoside, nucleotide, lipid, saturated or unsaturated
 CC fatty acid, diol, carbohydrate, aromatic compound, vitamin, cofactor,
 CC polyketide or enzyme. The amino acids produced can be lysine, glutamine,
 CC glutamate, alanine, aspartate, glycine, serine, threonine, methionine,
 CC cysteine, valine, leucine, isoleucine, arginine, proline, histidine,
 CC tyrosine, phenylalanine, or tryptophan. The fine chemical production can
 CC be modulated. The presence of (I) or HA proteins encoded by then are
 CC used for diagnosing the presence or activity of Corynebacterium
 CC diphtheriae. (I) can be used to map the C. glutamicum genome or can be
 CC used as markers for genetically engineered Corynebacterium or
 CC Brevibacterium. The HA proteins encoded by the (I) are used to maintain
 CC homeostasis in C. glutamicum or help the microorganism to adapt to
 CC different environmental conditions.
 XX

SQ Sequence 341 AA;

Query Match 90.5%; Score 19; DB 22; Length 341;
 Best Local Similarity 66.7%; Pred. No. 2.le+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 239 eagtss 244

RESULT 34

AA79144
 ID AAB79144 standard; Protein; 341 AA.

XX AC AAB79144;

XX 30-APR-2001 (first entry)

XX Corynebacterium glutamicum HA protein sequence SEQ ID NO:244.

XX Corynebacterium glutamicum; homeostasis; adaptation; HA protein;
 KW fine chemical production; organic acid; proteinogenic amino acid;
 KW nonproteinogenic amino acid; purine base; pyrimidine base; nucleoside;
 KW nucleotide; lipid; saturated fatty acid; unsaturated fatty acid; diol;
 KW carbohydrate; aromatic compound; vitamin; cofactor; polyketide; enzyme;
 KW diagnosis; Corynebacterium diphtheriae; genetic engineering;
 KW Brevibacterium; environmental condition.

XX OS Corynebacterium glutamicum.

XX WO200100842-A2.

XX 04-JAN-2001.

XX 23-JUN-2000; 2000WO-IB00911.

XX 25-JUN-1999; 99US-0141031.

XX 08-JUL-1999; 99DE-1031636.

XX 09-JUL-1999; 99DE-1032125.

XX 09-JUL-1999; 99DE-1032126.

XX 09-JUL-1999; 99DE-1032127.

XX 09-JUL-1999; 99DE-1032128.

XX 09-JUL-1999; 99DE-1032129.

XX 09-JUL-1999; 99DE-1032226.

XX 14-JUL-1999; 99DE-1032920.

XX 14-JUL-1999; 99DE-1032922.

XX 14-JUL-1999; 99DE-1032924.

XX 14-JUL-1999; 99DE-1032928.

XX 14-JUL-1999; 99DE-1032930.

XX 14-JUL-1999; 99DE-1032933.

XX 14-JUL-1999; 99DE-1032935.

XX 14-JUL-1999; 99DE-1032973.

XX 14-JUL-1999; 99DE-1033002.

XX 14-JUL-1999; 99DE-1033003.

XX 14-JUL-1999; 99DE-1033005.

XX 31-AUG-1999; 99DE-1033006.

XX 31-AUG-1999; 99DE-1041378.

XX 31-AUG-1999; 99DE-1041379.

XX 31-AUG-1999; 99DE-1041390.

XX 31-AUG-1999; 99DE-1041391.

XX 03-SEP-1999; 99DE-1042088.

XX (BADI) BASF AG.

XX Pompejus M, Kroeger B, Schroeder H, Zelder O, Haberhauer G;

XX WPI; 2001-061974/07.

XX N-PSDB; AAF71259.

XX New isolated Corynebacterium glutamicum nucleic acid for production or
 PT modulation of production of fine chemicals such as amino acids,

PT nucleosides, nucleotides, lipids, fatty acids, carbohydrates, vitamins
 or enzymes -

XX Claim 20; Page 466-467; 712pp; English.

XX AAF71138 to AAF71357 encode the Corynebacterium glutamicum homeostasis
 and adaptation (HA) proteins given in AAB79023 to AAB79242. The
 C. glutamicum HA genes (I) can be used in vectors for expression in host
 cells and production of fine chemicals, such as, an organic acid,
 CC proteinogenic or nonproteinogenic amino acid (preferred), purine or
 CC pyrimidine base, nucleoside, nucleotide, lipid, saturated or unsaturated
 CC fatty acid, diol, carbohydrate, aromatic compound, vitamin, cofactor,
 CC polyketide or enzyme. The amino acids produced can be lysine, glutamine,
 CC glutamate, alanine, aspartate, glycine, serine, threonine, methionine,
 CC cysteine, valine, leucine, isoleucine, arginine, proline, histidine,
 CC tyrosine, phenylalanine, or tryptophan. The fine chemical production can
 CC be modulated. The presence of (I) or HA proteins encoded by then are
 CC used for diagnosing the presence or activity of Corynebacterium
 CC diphtheriae. (I) can be used to map the C. glutamicum genome or can be
 CC used as markers for genetically engineered Corynebacterium or
 CC Brevibacterium. The HA proteins encoded by the (I) are used to maintain
 CC homeostasis in C. glutamicum or help the microorganism to adapt to
 CC different environmental conditions.

XX Sequence 341 AA;

Query Match 90.5%; Score 19; DB 22; Length 341;
 Best Local Similarity 66.7%; Pred. No. 2.le+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 239 eagtss 244

RESULT 35

AAW44368

ID AAW44368 standard; Protein; 349 AA.

XX AC AAW44368;

XX 29-MAY-1998 (first entry)

XX Aspergillus nidulans metallo-protease pepH.

XX Aspergillus nidulans; metallo-protease; pepH; protein degradation;
 KW fungus; food processing.

XX OS Aspergillus nidulans.

XX WO9746689-A1.

XX 11-DEC-1997.

XX 05-JUN-1997; 97WO-EP02982.

XX 05-JUN-1996; 96EP-0201579.

XX (KONN) GIST-BROCADES BV.

XX Van Den Hombergh JPTW, Visser J;

XX WPI; 1998-042197/04.

XX N-PSDB; AAV15305, AAV15306.

XX Metallo-protease deficient fungus with site selected DNA disruption
 PT - and Aspergillus metallo-protease genes, useful in protein
 XX production to reduce protease activity hence protein degradation
 PS Example 4; Page 26-27; 53pp; English.

XX The present sequence represents a metallo-protease, pepH, from

CC Aspergillus nidulans from the present invention. The present invention
 CC describes a new protease deficient filamentous fungus, optionally with
 CC reduced extracellular acid protease activity, containing a site selected
 CC disruption of DNA resulting in reduced metallo-protease activity. The
 CC fungi are useful for the production of (heterologous and homologous)
 CC proteins e.g. for food processing, since reduced protease activity
 CC minimises the chance that, and rate at which, the proteins are degraded
 CC during production. DNA sequences encoding metallo-proteases can be
 CC used to produce metallo-protease deficient fungi, by transforming a
 CC filamentous fungus mutant with the constructs and selecting a
 CC transformed fungus with reduced metallo-protease activity. They are also
 CC useful for producing filamentous fungal metallo-protease, by culturing
 CC filamentous fungi transformed with the constructs under suitable
 CC conditions for sequence expression and recovering the metallo-protease.
 CC Such metallo-proteases are useful to assess in vitro whether proteins
 CC which it is proposed to produce from a fungal host are susceptible to
 CC the protease, so determining which metallo-protease genes need to be
 CC inactivated in the host. They are also useful in industrial processes.
 CC
 CC Sequence 349 AA;

Query Match 90.5%; Score 19; DB 19; Length 349;
 Best Local Similarity 66.7%; Pred. No. 2.1e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 237 eagsts 242

RESULT 36
 AAR14147
 ID AAR14147 standard; Protein; 352 AA.

XX AC AAR14147;

XX DT 12-DEC-1991 (first entry)

XX DE Pre-pro neutral protease II.

XX KW recombinant enzyme.

XX OS Aspergillus oryzae.

XX FH Key Location/Qualifiers
 XX FT Peptide 176..352
 XX FT /note="mature neutral protease II"

XX PN JP03198779-A.

XX PD 29-AUG-1991.

XX PF 27-DEC-1989; 89JP-0336737.

XX PR 27-DEC-1989; 89JP-0336737.

XX PA (SHOK-) SHOKUHN SANGYO KOS.

XX WPI: 1991-299435/41.

XX DR N-PSDB; AAO13852.

XX PT Neutral protease II gene from Aspergillus - used to produce
 XX recombinant enzyme by expression in Saccharomyces.

XX PS Claim; Fig 4; 15pp; Japanese.

XX CC The pre-pro neutral protease II is derived from Aspergillus
 CC oryzae and is the precursor to neutral protease II. The neutral
 CC protease II can be expressed in Saccharomyces cerevisiae and can be
 CC produced efficiently and in a secreted form.

XX SQ Sequence 352 AA;

Query Match 90.5%; Score 19; DB 12; Length 352;
 Best Local Similarity 66.7%; Pred. No. 2.2e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 240 eagsts 245

RESULT 37

AAU36356

ID AAU36356 standard; Protein; 363 AA.

XX AC AAU36356;

XX DT 14-FEB-2002 (first entry)

XX DE Pseudomonas aeruginosa cellular proliferation protein #346.

XX KW Antisense: prokaryotic cellular proliferation protein;
 KW antibiotic; antibacterial; drug design.

XX OS Pseudomonas aeruginosa.

XX PN WO200170955-A2.

XX PD 27-SEP-2001.

XX PF 21-MAR-2001; 2001WO-US09180.

XX PR 21-MAR-2000; 2000US-191078P.

XX PR 23-MAY-2000; 2000US-206848P.

XX PR 26-MAY-2000; 2000US-207727P.

XX PR 23-OCT-2000; 2000US-242578P.

XX PR 27-NOV-2000; 2000US-253625P.

XX PR 22-DEC-2000; 2000US-257931P.

XX PR 16-FEB-2001; 2001US-269308P.

XX PA (ELIT-) ELITRA PHARM INC.

XX PI Haselbeck R, Ohlsen KL, Zyskind JW, Wall D, Trawick JD, Carr GJ;
 XX PI Yamamoto RT, Xu HH;

XX WPI: 2001-611495/70.

XX DR N-PSDB; AAS54215.

XX PT New polynucleotides for the identification and development of
 XX antibiotics, comprise sequences of antisense nucleic acids -

XX PS Example 3; Seq ID No 11949; 51pp; English.

XX CC The invention relates to antisense inhibitors of genes essential to
 CC prokaryotic cellular proliferation, their use in identifying the
 CC genes, their use in the discovery of novel antibiotics, the essential
 CC genes themselves and the encoded proteins. The prokaryotes used are
 CC Escherichia coli, Staphylococcus aureus, Salmonella typhi, Klebsiella
 CC pneumoniae, Pseudomonas aeruginosa and Enterococcus faecalis. The
 CC invention is also useful for the identification of potential new targets
 CC for antibiotic development. The antisense nucleic acids can also be used
 CC to identify proteins used in proliferation, to express these proteins,
 CC and to obtain antibodies capable of binding to the expressed proteins.
 CC The proteins can be used to screen compounds in rational drug discovery
 CC programmes. The antisense nucleic acid sequence is also useful to screen
 CC for homologous nucleic acids which are required for cell proliferation in
 CC a wide variety of organisms. The present sequence represents an
 CC essential prokaryotic cellular proliferation protein.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

SQ Sequence 363 AA;

Query Match 90.5%; Score 19; DB 22; Length 363;
Best Local Similarity 66.7%; Pred. No. 2.2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 eagxxs 6
||| |
Db 196 eagass 201

RESULT 38

AAAY05663
ID AAY05663 standard; Protein; 366 AA.

XX AC AAY05663;

XX DT 19-JUL-1999 (first entry)

XX DE Maize caffeic O-methyltransferase.

XX KW Maize; corn; caffeic O-methyltransferase; lignin; transgenic plant.

XX OS Zea mays.

XX PN WO9910498-A2.

XX PD 04-MAR-1999.

XX PF 24-AUG-1998; 98WO-US17519.

XX PR 12-MAY-1998; 98US-0076851.

XX PR 27-AUG-1997; 97US-0057082.

XX PA (PION-) PIONEER HI-BRED INT INC.

XX PI Bowen BA, Helentjaris TG, Wang X;

XX DR WPI; 1999-204667/17.

XX DR N-PSDB; AAX25202.

XX PT Nucleic acids encoding plant lignin biosynthesis enzymes - used to
transform plants to modulate lignin biosynthesis

XX PS Claim 9; Page 96-97; 166pp; English.

XX CC The present sequence is a caffeic O-methyltransferase of maize,
encoded by a clone (see AAX25202) isolated from a maize line B73 cDNA
library. The invention provides methods and compositions relating
to altering lignin biosynthesis and/or the lignin composition of
plants. Isolated nucleic acids (see AAX25196-216) that code for
proteins (see AAY05657-77) involved in lignin biosynthesis are
claimed. Also claimed are recombinant expression cassettes, host
cells (especially maize or sorghum), and transgenic plants and
seeds. The claimed nucleic acids can be used to transform a plant
to modulate lignin biosynthesis. A claimed method involves
transforming a plant cell with a recombinant expression cassette
comprising a lignin biosynthesis polynucleotide operably linked to
a promoter, growing the plant cell under plant growing conditions,
and inducing expression of the polynucleotide for a time sufficient
to modulate (preferably increase) lignin biosynthesis in the plant.
The plant lignins can be used as chemical feedstock. Plant
material of increased lignin content can be used as a fuel source,
and in the pulp and paper industry. Decreased lignin content
improves the digestibility of fodder crops.

XX SQ Sequence 366 AA;

Query Match 90.5%; Score 19; DB 20; Length 366;
Best Local Similarity 66.7%; Pred. No. 2.2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 eagxxs 6
||| |
Db 115 eagtas 120

RESULT 39

AAG20561

ID AAG20561 standard; Protein; 377 AA.

XX AC AAG20561;

XX DT 17-OCT-2000 (first entry)

XX DE Arabidopsis thaliana protein fragment SEQ ID NO: 22802.

XX KW Protein identification; signal transduction pathway; metabolic pathway;
XX KW hybridisation assay; genetic mapping; gene expression control; promoter;
XX KW termination sequence.

XX OS Arabidopsis thaliana.

XX PN EPI033405-A2.

XX PD 06-SEP-2000.

XX PF 25-FEB-2000; 2000EP-0301439.

XX PR 25-FEB-1999; 99US-0121825.

XX PR 05-MAR-1999; 99US-0123180.

XX PR 09-MAR-1999; 99US-0123548.

XX PR 23-MAR-1999; 99US-0125788.

XX PR 25-MAR-1999; 99US-0126264.

XX PR 29-MAR-1999; 99US-0126785.

XX PR 01-APR-1999; 99US-0127462.

XX PR 06-APR-1999; 99US-0128234.

XX PR 08-APR-1999; 99US-0128714.

XX PR 16-APR-1999; 99US-0129845.

XX PR 19-APR-1999; 99US-0130077.

XX PR 21-APR-1999; 99US-0130449.

XX PR 23-APR-1999; 99US-0130510.

XX PR 28-APR-1999; 99US-0130891.

XX PR 30-APR-1999; 99US-0131449.

XX PR 30-APR-1999; 99US-0132048.

XX PR 04-MAY-1999; 99US-0132407.

XX PR 05-MAY-1999; 99US-0132485.

XX PR 06-MAY-1999; 99US-0132486.

XX PR 07-MAY-1999; 99US-0132487.

XX PR 11-MAY-1999; 99US-0132863.

XX PR 14-MAY-1999; 99US-0134256.

XX PR 14-MAY-1999; 99US-0134218.

XX PR 14-MAY-1999; 99US-0134219.

XX PR 14-MAY-1999; 99US-0134221.

XX PR 14-MAY-1999; 99US-0134370.

XX PR 18-MAY-1999; 99US-0134768.

XX PR 19-MAY-1999; 99US-0134941.

XX PR 20-MAY-1999; 99US-0135124.

XX PR 21-MAY-1999; 99US-0135353.

XX PR 24-MAY-1999; 99US-0135629.

XX PR 25-MAY-1999; 99US-0136021.

XX PR 27-MAY-1999; 99US-0136392.

XX PR 28-MAY-1999; 99US-0136782.

XX PR 01-JUN-1999; 99US-0137222.

XX PR 03-JUN-1999; 99US-0137528.

XX PR 04-JUN-1999; 99US-0137502.

XX PR 07-JUN-1999; 99US-0137724.

XX PR 08-JUN-1999; 99US-0138094.

XX PR 10-JUN-1999; 99US-0138540.

XX PR 10-JUN-1999; 99US-0138847.

XX PR 14-JUN-1999; 99US-0139119.

XX PR 16-JUN-1999; 99US-0139452.

XX PR 16-JUN-1999; 99US-0139453.

PR 17-JUN-1999; 99US-0139492.
PR 18-JUN-1999; 99US-0139454.
PR 18-JUN-1999; 99US-0139455.
PR 18-JUN-1999; 99US-0139456.
PR 18-JUN-1999; 99US-0139457.
PR 18-JUN-1999; 99US-0139458.
PR 18-JUN-1999; 99US-0139459.
PR 18-JUN-1999; 99US-0139460.
PR 18-JUN-1999; 99US-0139461.
PR 18-JUN-1999; 99US-0139462.
PR 18-JUN-1999; 99US-0139463.
PR 18-JUN-1999; 99US-0139750.
PR 18-JUN-1999; 99US-0139763.
PR 21-JUN-1999; 99US-0139817.
PR 22-JUN-1999; 99US-0139899.
PR 23-JUN-1999; 99US-0140353.
PR 23-JUN-1999; 99US-0140354.
PR 24-JUN-1999; 99US-0140695.
PR 28-JUN-1999; 99US-0140823.
PR 29-JUN-1999; 99US-0140991.
PR 30-JUN-1999; 99US-0141287.
PR 01-JUL-1999; 99US-0141842.
PR 01-JUL-1999; 99US-0142154.
PR 02-JUL-1999; 99US-0142055.
PR 06-JUL-1999; 99US-0142390.
PR 08-JUL-1999; 99US-0142803.
PR 09-JUL-1999; 99US-0142920.
PR 12-JUL-1999; 99US-0142977.
PR 13-JUL-1999; 99US-0143542.
PR 14-JUL-1999; 99US-0143624.
PR 15-JUL-1999; 99US-0144005.
PR 16-JUL-1999; 99US-0144085.
PR 16-JUL-1999; 99US-0144086.
PR 19-JUL-1999; 99US-0144325.
PR 19-JUL-1999; 99US-0144331.
PR 19-JUL-1999; 99US-0144332.
PR 19-JUL-1999; 99US-0144333.
PR 19-JUL-1999; 99US-0144334.
PR 19-JUL-1999; 99US-0144335.
PR 20-JUL-1999; 99US-0144352.
PR 20-JUL-1999; 99US-0144632.
PR 20-JUL-1999; 99US-0144884.
PR 21-JUL-1999; 99US-0144814.
PR 21-JUL-1999; 99US-0145086.
PR 21-JUL-1999; 99US-0145088.
PR 22-JUL-1999; 99US-0145085.
PR 22-JUL-1999; 99US-0145087.
PR 22-JUL-1999; 99US-0145089.
PR 22-JUL-1999; 99US-0145192.
PR 23-JUL-1999; 99US-0145145.
PR 23-JUL-1999; 99US-0145218.
PR 23-JUL-1999; 99US-0145224.
PR 26-JUL-1999; 99US-0145276.
PR 27-JUL-1999; 99US-0145913.
PR 27-JUL-1999; 99US-0145918.
PR 27-JUL-1999; 99US-0145919.
PR 28-JUL-1999; 99US-0145951.
PR 02-AUG-1999; 99US-0146386.
PR 02-AUG-1999; 99US-0146388.
PR 02-AUG-1999; 99US-0146389.
PR 03-AUG-1999; 99US-0147038.
PR 04-AUG-1999; 99US-0147204.
PR 04-AUG-1999; 99US-0147302.
PR 05-AUG-1999; 99US-0147192.
PR 05-AUG-1999; 99US-0147260.
PR 06-AUG-1999; 99US-0147303.
PR 06-AUG-1999; 99US-0147416.
PR 09-AUG-1999; 99US-0147935.
PR 09-AUG-1999; 99US-0147935.
PR 10-AUG-1999; 99US-0148171.
PR 11-AUG-1999; 99US-0148319.
PR 12-AUG-1999; 99US-0148341.
PR 13-AUG-1999; 99US-0148565.

PR 13-AUG-1999; 99US-0148684.
PR 16-AUG-1999; 99US-0149368.
PR 17-AUG-1999; 99US-0149175.
PR 18-AUG-1999; 99US-0149426.
PR 20-AUG-1999; 99US-0149722.
PR 20-AUG-1999; 99US-0149723.
PR 20-AUG-1999; 99US-0149929.
PR 23-AUG-1999; 99US-0149902.
PR 23-AUG-1999; 99US-0149930.
PR 25-AUG-1999; 99US-0150566.
PR 26-AUG-1999; 99US-0150884.
PR 27-AUG-1999; 99US-0151065.
PR 27-AUG-1999; 99US-0151066.
PR 27-AUG-1999; 99US-0151080.
PR 30-AUG-1999; 99US-0151303.
PR 31-AUG-1999; 99US-0151438.
PR 01-SEP-1999; 99US-0151930.
PR 07-SEP-1999; 99US-0152363.
PR 10-SEP-1999; 99US-0153070.
PR 13-SEP-1999; 99US-0153758.
PR 15-SEP-1999; 99US-0154018.
PR 16-SEP-1999; 99US-0154039.
PR 20-SEP-1999; 99US-0154779.
PR 22-SEP-1999; 99US-0155139.
PR 23-SEP-1999; 99US-0155486.
PR 24-SEP-1999; 99US-0156559.
PR 28-SEP-1999; 99US-0156458.
PR 29-SEP-1999; 99US-0156596.
PR 04-OCT-1999; 99US-0157117.
PR 06-OCT-1999; 99US-0157753.
PR 06-OCT-1999; 99US-0157865.
PR 07-OCT-1999; 99US-0158029.
PR 08-OCT-1999; 99US-0158232.
PR 12-OCT-1999; 99US-0158369.
PR 13-OCT-1999; 99US-0159293.
PR 13-OCT-1999; 99US-0159294.
PR 13-OCT-1999; 99US-0159295.
PR 14-OCT-1999; 99US-0159329.
PR 14-OCT-1999; 99US-0159330.
PR 14-OCT-1999; 99US-0159331.
PR 14-OCT-1999; 99US-0159637.
PR 14-OCT-1999; 99US-0159638.
PR 18-OCT-1999; 99US-0159584.
PR 21-OCT-1999; 99US-0160741.
PR 21-OCT-1999; 99US-0160767.
PR 21-OCT-1999; 99US-0160768.
PR 21-OCT-1999; 99US-0160770.
PR 21-OCT-1999; 99US-0160814.
PR 21-OCT-1999; 99US-0160815.
PR 22-OCT-1999; 99US-0160980.
PR 22-OCT-1999; 99US-0160981.
PR 22-OCT-1999; 99US-0160989.
PR 25-OCT-1999; 99US-0161404.
PR 25-OCT-1999; 99US-0161405.
PR 25-OCT-1999; 99US-0161406.
PR 26-OCT-1999; 99US-0161359.
PR 26-OCT-1999; 99US-0161360.
PR 26-OCT-1999; 99US-0161361.
PR 28-OCT-1999; 99US-0161920.
PR 28-OCT-1999; 99US-0161992.
PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

Query Match 90.5%; Score 19; DB 21; Length 377;
Best Local Similarity 66.7%; Pred. No. 2.3e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaggxs 6
||| |
Db 75 eaggss 80

RESULT 40

AAG41800
ID AAG41800 standard; Protein; 377 AA.
XX
AC AAG41800;
XX
XX
DT 18-OCT-2000 (first entry)
XX
DE Arabidopsis thaliana protein fragment SEQ ID NO: 52052.
XX
KW Protein identification; signal transduction pathway; metabolic pathway;
KW hybridisation assay; genetic mapping; gene expression control; promoter;
KW termination sequence.
XX
OS Arabidopsis thaliana.
XX
XX EPI033405-A2.
XX
XX 06-SEP-2000.
XX
PF 25-FEB-2000; 2000EP-0301439.
XX
XX 25-FEB-1999; 99US-0121825.
XX 05-MAR-1999; 99US-0123180.
XX 09-MAR-1999; 99US-0123548.
XX 23-MAR-1999; 99US-0125788.
XX 25-MAR-1999; 99US-0126264.
XX 29-MAR-1999; 99US-0126785.
XX 01-APR-1999; 99US-0127462.
XX 06-APR-1999; 99US-0128234.
XX 08-APR-1999; 99US-0128714.
XX 16-APR-1999; 99US-0129845.
XX 19-APR-1999; 99US-0130077.
XX 21-APR-1999; 99US-0130449.
XX 23-APR-1999; 99US-0130510.
XX 23-APR-1999; 99US-0130891.
XX 28-APR-1999; 99US-0131449.
XX 30-APR-1999; 99US-0132048.
XX 30-APR-1999; 99US-0132407.
XX 04-MAY-1999; 99US-0132484.
XX 05-MAY-1999; 99US-0132485.
XX 06-MAY-1999; 99US-0132486.
XX 06-MAY-1999; 99US-0132487.
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XX 11-MAY-1999; 99US-0134256.
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XX 14-MAY-1999; 99US-0134370.
XX 18-MAY-1999; 99US-0134768.
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XX 20-MAY-1999; 99US-0135124.
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XX 24-MAY-1999; 99US-0135629.
XX 25-MAY-1999; 99US-0136021.
XX 27-MAY-1999; 99US-0136392.
XX 28-MAY-1999; 99US-0136782.
XX 01-JUN-1999; 99US-0137222.
XX 03-JUN-1999; 99US-0137528.
XX 04-JUN-1999; 99US-0137502.
XX 07-JUN-1999; 99US-0137724.
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PR 23-AUG-1999; 99US-0149902.
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PR 26-AUG-1999; 99US-0150884.
PR 27-AUG-1999; 99US-0151065.
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PR 27-AUG-1999; 99US-0151080.
PR 30-AUG-1999; 99US-0151303.
PR 31-AUG-1999; 99US-0151438.
PR 01-SEP-1999; 99US-0151930.
PR 07-SEP-1999; 99US-0152363.
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PR 28-OCT-1999; 99US-0161992.
PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

Query Match 90.5%; Score 19; DB 21; Length 377;
Best Local Similarity 66.7%; Pred. No. 2.3e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 eagxxs 6
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Db 75 eagsss 80

RESULT 41
AAG24042
ID AAG24042 standard; Protein; 386 AA.
XX AC AAG24042;
XX DT 17-Oct-2000 (first entry)
XX

DE Arabidopsis thaliana protein fragment SEQ ID NO: 27567.
XX
KW Protein identification; signal transduction pathway; metabolic pathway;
KW hybridisation assay; genetic mapping; gene expression control; promoter;
XX termination sequence.
OS Arabidopsis thaliana.
XX
PN EP1033405-A2.
XX
PD 06-SEP-2000.
XX
PF 25-FEB-2000; 2000EP-0301439.
XX
PR 25-FEB-1999; 99US-0121825.
PR 05-MAR-1999; 99US-0123180.
PR 03-MAR-1999; 99US-0123548.
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PR 27-AUG-1999; 99US-0151080.

PR 30-AUG-1999; 99US-0151303.
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PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

Query Match 90.5%; Score 19; DB 21; Length 386;
Best Local Similarity 66.7%; Pred. NO. 2.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
Db 46 eagaas 51

RESULT 42
AAP70581
ID AAP70581 standard; Protein; 390 AA.
XX AAP70581;
XX AC
XX DT 03-MAY-1991 (first entry)
XX DE Protease biosynthetic protein.
XX DE Saccharomycosis; yeast.
XX KW JP62104578-A.
XX PN
XX PD 15-MAY-1987.

XX 31-OCT-1985; 85JP-0244893.
 XX
 PR 31-OCT-1985; 85JP-0244893.
 XX
 PA (FUKU/) FUKUI S.
 XX
 DR WPI; 1987-173695/25.
 DR N-PSDB; AAN70927.
 XX
 PT Protease prodn. - by culturing microorganism transformed with
 PT vector derived from saccharomycosis.
 XX
 PS Disclosure; Fig 1; 9pp; Japanese.
 XX
 CC Product is a biosynthetic component involved in the synthesis of
 CC protease. The protein may be produced from a transformed *S.cerevisiae*
 CC expression system for the large scale production of protease.
 XX
 SQ Sequence 390 AA;
 XX
 Query Match 90.5%; Score 19; DB 8; Length 390;
 Best Local Similarity 66.7%; Pred. No. 2.4e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 eagxss 6
 Db 266 eagsss 271
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 RESULT 43
 ID AAY35147
 ID AAY35147 standard; Protein; 393 AA.
 XX
 AC AAY35147;
 XX
 DT 13-SEP-1999 (first entry)
 XX
 DE Chlamydia pneumoniae transmembrane protein sequence.
 XX
 KW Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;
 KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;
 KW vaccine; neutralising epitope.
 XX
 OS Chlamydia pneumoniae.
 XX
 PN WO9927105-A2.
 XX
 PD 03-JUN-1999.
 XX
 PF 20-NOV-1998; 98WO-IB01890.
 XX
 PR 04-NOV-1998; 98US-0107078.
 PR 21-NOV-1997; 97FR-0014673.
 XX
 PA (GEST) GENSET.
 XX
 PI Griffois R;
 XX
 DR WPI; 1999-357842/30.
 XX
 PT Genome sequence of Chlamydia pneumoniae
 XX
 PS Page 1016; Disclosure; 1912pp; English.
 XX
 CC AAY34584-Y35879 represent the proteins encoded by all the open reading
 CC frames in the complete genome (see AAX91990) of Chlamydia pneumoniae.
 CC C. pneumoniae causes respiratory disease such as pneumonia and
 CC bronchitis and is thought to be a contributing factor in heart
 CC disease, sarcoidosis, sinusitis, purulent otitis media, erythema
 CC nodosum or pharyngitis. The polypeptides encoded by the open reading
 CC frames of the C. pneumoniae genome (see AAY34584-Y35879) can be used in

CC immunogenic compositions as vaccines. Vectors containing C. pneumoniae
 CC nucleotides sequences can also be used as immunogenic compositions,
 CC especially where the vector directs the expression of a neutralising
 CC epitope of C. pneumoniae.
 XX
 SQ Sequence 393 AA;
 XX
 Query Match 90.5%; Score 19; DB 20; Length 393;
 Best Local Similarity 66.7%; Pred. No. 2.4e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 eagxss 6
 Db 101 eagsss 106
 ||| |
 RESULT 44
 ID AAY28643
 ID AAY28643 standard; Protein; 422 AA.
 XX
 AC AAY28643;
 XX
 DT 03-NOV-1999 (first entry)
 XX
 DE Human serine protease inhibitor from cDNA clone HETDK50.
 XX
 KW Human serine protease inhibitor from cDNA clone HETDK50; fusion protein;
 KW serpin; serine protease; human pre-alpha-1-antitrypsin precursor;
 KW extracellular matrix degradation; multiple sclerosis; cancer; arthritis;
 KW inflammation; immune system disorder; neurodegenerative disorder;
 KW Kallmann's syndrome; Down's syndrome; Alzheimer's; secreted protein;
 KW galactorrhea; hypogonadism; somatostatin; protein purification.
 XX
 OS Homo sapiens.
 XX
 FH Key
 FT Peptide 1..19
 FT /label= Signal_peptide
 FT Protein 20..422
 FT /label= Mature_serine_protease_inhibitor
 XX
 PN WO9940183-A1.
 XX
 PD 12-AUG-1999.
 XX
 PF 04-FEB-1999; 99WO-US02292.
 XX
 PR 06-FEB-1998; 98US-0073961.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Ni J, Ruben SM;
 XX
 DR WPI; 1999-508502/42.
 DR N-PSDB; AAX80907.
 XX
 PT New isolated human serine protease and serpin polypeptides, used to
 PT develop products for treating e.g. immune disorders, cancers,
 PT inflammation, transplant rejection or infections, or as food
 PT additives
 XX
 PS Claim 11; Pages 83-85; 99pp; English.
 XX
 CC The present sequence is a serine protease inhibitor (serpin) from cDNA
 CC clone HETDK50 which is obtained from human endometrial tumour tissue
 CC cDNA library. The protein shows a high degree of sequence similarity
 CC human pre-alpha-1-antitrypsin precursor. The serpin and its
 CC coding sequence are used in the diagnosis and treatment of disorders
 CC related to abnormal level of the protein or mutation in the nucleotide
 CC sequence. The serpin can be used for treating disorders characterised by
 CC degradation of extracellular matrix, e.g. cancer, arthritis, multiple
 CC sclerosis and immune system disorders, for treating wasting associated

CC with excessive protease production during inflammation or
 CC neurodegenerative disorders e.g. Kallmann's and Down's syndromes,
 CC Alzheimer's and Huntington's diseases. It may also be used to reduce
 CC excess levels of prolactin in the treatment of galactorrhoea and
 CC hypogonadism, and decrease the amount of free circulating somatostatin to
 CC prevent somatostatin's inhibitory effect on the release of growth
 CC hormone. The fusion of this protein to His-tag, HA-tag, IgG domains,
 CC etc. facilitates protein purification and fusion to IgG-1, IgG-3 and
 CC albumin increases the half life time in vivo.

XX
 SQ Sequence 422 AA;

Query Match 90.5%; Score 19; DB 20; Length 422;

Best Local Similarity 66.7%; Pred. No. 2.6e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

||| |

Db 370 eaggaas 375

RESULT 45

AAB74691

ID AAB74691 standard; Protein: 422 AA.

XX AC AAB74691;

XX 12-JUN-2001 (first entry).

XX Human protease and protease inhibitor PPIM-24.

XX Human; protease; protease inhibitor; protease and protease inhibitor;
 KW PPIM; identification; diagnosis; anti-human immunodeficiency virus; HIV;
 KW antidiabetic; immunostimulant; immunomodulator; antiinflammatory;
 KW antithyroid; immunosuppressive; nephrotropic; antigout; thyromimetic;
 KW cytostatic; antibacterial; fungicide; protozoacide; antiarteriosclerotic;
 KW antiatherosclerotic; antipsoriatic; virucide; hepatotropic; gene therapy;
 KW autoimmune disorder; inflammatory disease; AIDS; Digeorge's syndrome;
 KW severe combined immunodeficiency disease; SCID; Chediak-Higashi syndrome;
 KW Cushing's disease; Addison's disease; autoimmune thyroiditis; gout;
 KW Crohn's disease; diabetes mellitus; Good pasture's syndrome; infection;
 KW Grave's diseases; Hashimoto's thyroiditis; Sjogren's syndrome; cancer;
 KW Werner's syndrome; cell proliferative disorder; arteriosclerosis;
 KW atherosclerosis; cirrhosis; hepatitis; psoriasis.

XX Homo sapiens.

XX W0200110903-A2.

XX 15-FEB-2001.

XX 09-AUG-2000; 2000WO-US21878.

XX 09-AUG-1999; 99US-0147986.

XX 21-OCT-1999; 99US-0160807.

XX (INCY-) INCYTE GENOMICS INC.

XX Yue H, Lal P, Tang YT, Bandman O, Baughn MR, Azimzai Y, Lu DAM;

PI Yang J;

XX WPI; 2001-202760/20.

XX N-PSDB; AAF81737.

XX New protease (inhibitors) useful for diagnosis and treatment of
 PT autoimmune/inflammatory disorders such as acquired immunodeficiency
 PT syndrome, Cushing's disease, Addison's disease and cell proliferative
 PT disorders such as cancer -

PS Claim 1; Page 112-113; 134pp; English.

XX AAF81714 to AAF81740 encode the human proteases and protease inhibitors

CC (PPIMs) given in AAB74668 to AAB74694. The PPIMs can have activities such
 CC as: anti-human immunodeficiency virus (HIV); antidiabetic; antithyroid;
 CC immunostimulant; immunomodulator; antiinflammatory; immunosuppressive;
 CC nephrotropic; antigout; thyromimetic; cytostatic; antibacterial;
 CC fungicide; protozoacide; antiarteriosclerotic; antiatherosclerotic;
 CC virucide; antipsoriatic; and hepatotropic. PPIM polynucleotide and
 CC protein sequences can be used in the diagnosis, treatment and prevention
 CC of autoimmune/inflammatory disorders such as AIDS, Digeorge's syndrome,
 CC severe combined immunodeficiency disease (SCID), Chediak-Higashi
 CC syndrome, Cushing's disease, Addison's disease, autoimmune thyroiditis,
 CC Crohn's disease, diabetes mellitus, Good pasture's syndrome, gout,
 CC Grave's diseases, Hashimoto's thyroiditis, Sjogren's syndrome, Werner's
 CC syndrome, viral, bacterial, fungal, parasitic, protozoal, and helminthic
 CC infections and cell proliferative disorder such as arteriosclerosis,
 CC atherosclerosis, cirrhosis, hepatitis, psoriasis and cancer. PPIM
 CC polynucleotide sequences can be used in somatic or germline gene therapy
 CC and in diagnosis of diseases. They can also be used in generating
 CC hybridisation probes useful in mapping the naturally occurring genomic
 CC sequences and in molecular biology techniques.

XX
 SQ Sequence 422 AA;

Query Match 90.5%; Score 19; DB 22; Length 422;

Best Local Similarity 66.7%; Pred. No. 2.6e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

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Db 370 eaggaas 375

RESULT 46

AAG41388

ID AAG41388 standard; Protein: 427 AA.

XX AC AAG41388;

XX 18-OCT-2000 (first entry)

XX Arabidopsis thaliana protein fragment SEQ ID NO: 51488.

XX Protein identification; signal transduction pathway; metabolic pathway;
 KW hybridisation assay; genetic mapping; gene expression control; promoter;
 KW termination sequence.

XX Arabidopsis thaliana.

XX EP1033405-A2.

XX 06-SEP-2000.

XX 25-FEB-2000; 2000EP-0301439.

XX 25-FEB-1999; 99US-0121825.

XX 05-MAR-1999; 99US-0123180.

XX 23-MAR-1999; 99US-0123548.

XX 23-MAR-1999; 99US-0125788.

XX 25-MAR-1999; 99US-0126264.

XX 29-MAR-1999; 99US-0126785.

XX 01-APR-1999; 99US-0127462.

XX 06-APR-1999; 99US-0128234.

XX 08-APR-1999; 99US-0128714.

XX 16-APR-1999; 99US-0129845.

XX 19-APR-1999; 99US-0130077.

XX 21-APR-1999; 99US-0130449.

XX 23-APR-1999; 99US-0130510.

XX 28-APR-1999; 99US-0130891.

XX 30-APR-1999; 99US-0131449.

XX 30-APR-1999; 99US-0132048.

XX 04-MAY-1999; 99US-0132407.

XX 05-MAY-1999; 99US-0132484.

XX 05-MAY-1999; 99US-0132485.

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PR 28-MAY-1999; 99US-0136782.
PR 01-JUN-1999; 99US-0137222.
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PR 27-AUG-1999; 99US-0151080.
PR 30-AUG-1999; 99US-0151303.
PR 31-AUG-1999; 99US-0151438.
PR 01-SEP-1999; 99US-0151930.
PR 07-SEP-1999; 99US-0152363.
PR 10-SEP-1999; 99US-0153070.
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Query Match 90.58; Score 19; DB 21; Length 427;
Best Local Similarity 66.78; Pred. No. 2.6e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
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Db 6 eagsss 11

RESULT 47
AAG41387
ID AAG41387 standard; Protein; 431 AA.

XX AC AAG41387;

XX DT 18-OCT-2000 (first entry)

XX DE Arabidopsis thaliana protein fragment SEQ ID NO: 51487.

XX KW Protein identification; signal transduction pathway; metabolic pathway;
KW hydriisation assay; genetic mapping; gene expression control; promoter;
KW termination sequence.

XX OS Arabidopsis thaliana.

XX PN EPI033405-A2.

XX PD 06-SEP-2000.

XX PF 25-FEB-2000; 2000EP-0301439.

XX PR 25-FEB-1999; 99US-0121825.

PR 05-MAR-1999; 99US-0123180.

PR 09-MAR-1999; 99US-0123548.

PR 23-MAR-1999; 99US-0125788.

PR 25-MAR-1999; 99US-0126264.

PR 29-MAR-1999; 99US-0126785.

PR 01-APR-1999; 99US-0127462.

PR 06-APR-1999; 99US-0128234.

PR 08-APR-1999; 99US-0128714.

PR 16-APR-1999; 99US-0129845.

PR 19-APR-1999; 99US-0130077.

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PR 28-APR-1999; 99US-0130891.

PR 30-APR-1999; 99US-0131449.

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PR 04-MAY-1999; 99US-0132407.

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PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

Query Match 90.5%; Score 19; DB 21; Length 431;
Best Local Similarity 66.7%; Pred. No. 2.6e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
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Db 10 eagsss 15

RESULT 48
AAG24041
ID AAG24041 standard; Protein; 435 AA.
XX AC AAG24041;
XX DT 17-OCT-2000 (first entry)
XX DE Arabidopsis thaliana protein fragment SEQ ID NO: 27566.
XX KW Protein identification; signal transduction pathway; metabolic pathway;
KW hybridisation assay; genetic mapping; gene expression control; promoter;
KW termination sequence.
XX OS Arabidopsis thaliana.
XX PN EF1033405-A2.
XX PD 06-SEP-2000.
XX PF 25-FEB-2000; 2000EP-0301439.
XX PR 25-FEB-1999; 99US-0121825.
PR 05-MAR-1999; 99US-0123180.
PR 09-MAR-1999; 99US-0123548.
PR 23-MAR-1999; 99US-0125788.
PR 25-MAR-1999; 99US-0126264.
PR 29-MAR-1999; 99US-0126785.
PR 01-APR-1999; 99US-0127462.
PR 06-APR-1999; 99US-0128234.
PR 08-APR-1999; 99US-0128714.
PR 16-APR-1999; 99US-0129845.
PR 19-APR-1999; 99US-0130077.
PR 21-APR-1999; 99US-0130449.
PR 23-APR-1999; 99US-0130510.
PR 28-APR-1999; 99US-0130891.
PR 30-APR-1999; 99US-0131449.
PR 30-APR-1999; 99US-0132048.
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PR 28-MAY-1999; 99US-0136782.
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PR 16-JUN-1999; 99US-0139453.
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PR 01-JUL-1999; 99US-0141842.
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PR 28-OCT-1999; 99US-0161993.

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PR 09-AUG-1999; 99US-0147935.
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 PR 25-AUG-1999; 99US-0150566.
 PR 26-AUG-1999; 99US-0150884.
 PR 27-AUG-1999; 99US-0151065.
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 PR 30-AUG-1999; 99US-0151303.
 PR 31-AUG-1999; 99US-0151438.
 PR 01-SEP-1999; 99US-0151930.
 PR 07-SEP-1999; 99US-0152363.
 PR 10-SEP-1999; 99US-0153070.
 PR 13-SEP-1999; 99US-0153758.
 PR 15-SEP-1999; 99US-0154018.
 PR 16-SEP-1999; 99US-0154039.
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 PR 05-OCT-1999; 99US-0157753.
 PR 06-OCT-1999; 99US-0157865.
 PR 07-OCT-1999; 99US-0158029.
 PR 08-OCT-1999; 99US-0158232.
 PR 12-OCT-1999; 99US-0158369.
 PR 13-OCT-1999; 99US-0159293.
 PR 13-OCT-1999; 99US-0159294.
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 PR 14-OCT-1999; 99US-0159331.
 PR 14-OCT-1999; 99US-0159637.
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 PR 21-OCT-1999; 99US-0160814.
 PR 21-OCT-1999; 99US-0160815.
 PR 22-OCT-1999; 99US-0160880.
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 PR 25-OCT-1999; 99US-0161404.
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 PR 26-OCT-1999; 99US-0161360.
 PR 26-OCT-1999; 99US-0161361.
 PR 28-OCT-1999; 99US-0161920.
 PR 28-OCT-1999; 99US-0161992.
 PR 28-OCT-1999; 99US-0161993.
 PR 29-OCT-1999; 99US-0162142.

Query Match 90.5%; Score 19; DB 21; Length 440;
 Best Local Similarity 66.7%; Pred. No. 2.7e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

Db 100 eagaas 105
 RESULT 50
 AAR31955
 ID AAR31955 standard; Protein; 441 AA.
 XX AC AAR31955;
 XX DT 06-JUN-1993 (first entry)
 XX DE Sequence encoded by glycoprotein G gene.
 XX KW IBR glycoprotein E gene; unique short 2 gene.
 XX OS Infectious bovine rhinotracheitis.
 XX PN WO9302104-A.
 XX PD 04-FEB-1993.
 XX PF 20-JUL-1992; 92WO-US06034.
 XX PR 18-JUL-1991; 91US-0732584.
 XX (SYTR) SYNTRO CORP.
 XX PI Cochran MD, Macdonald RD;
 XX DR WPI; 1993-058725/07.
 XX DR N-PSDB; AAQ36768.
 XX PT Recombinant infectious bovine rhinotracheitis virus - provides
 PT isolated DNA encoding gpE glyco:protein, gp6 glyco:protein and
 PT unique short 2 genes of the virus
 XX Example; Fig 8; 240pp; English.
 XX CC The sequence of approximately 1400 base pairs of the HindIII K
 CC fragment, starting approximately 2800 base pairs downstream of the
 CC HindIII K/HindIII O junction, are shown. The glycoprotein G (gpG)
 CC gene is transcribed away from the HindIII K/HindIII O junction.
 XX SQ Sequence 441 AA;
 Query Match 90.5%; Score 19; DB 14; Length 441;
 Best Local Similarity 66.7%; Pred. No. 2.7e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 eagxxs 6
 Db 265 eagaas 270
 Search completed: August 30, 2002, 15:05:43
 Job time: 6419 sec

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; INFORMATION FOR SEQ ID NO: 345:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 47 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: Protein
US-08-936-165A-345

Query Match 90.5%; Score 19; DB 4; Length 47;
Best Local Similarity 66.7%; Pred. No. 93;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |

Db 31 EAGATS 36

RESULT 2
US-08-856-253-2
; Sequence 2, Application US/08856253
; Patent No. 6288214
; GENERAL INFORMATION:
; APPLICANT: Hook, Magnus
; APPLICANT: Patti, Joseph M.
; APPLICANT: House-Pompeo, Karen
; APPLICANT: Sthanam, Narayana
; APPLICANT: Symersky, Jindrich
; TITLE OF INVENTION: COLLAGEN BINDING PROTEIN COMPOSITIONS
; TITLE OF INVENTION: AND METHODS OF USE
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Arnold, White & Durkee
; STREET: P.O. Box 4433
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.
; ZIP: 77210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/856,253
; FILING DATE: Concurrently Herewith
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/017,678
; FILING DATE: 16-MAY-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Kitchell, Barbara S.
; REGISTRATION NUMBER: 33,928
; REFERENCE/DOCKET NUMBER: TAMK:193
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (512) 474-7577
; TELEFAX: (512) 418-3000
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 159 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
US-08-856-253-2

Query Match 90.58; Score 19; DB 4; Length 159;
Best Local Similarity 66.7%; Pred. No. 3.le+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
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Db 28 EAGTSS 33
RESULT 3
US-08-856-253-4
; Sequence 4, Application US/08856253
; Patent No. 6288214
; GENERAL INFORMATION:
; APPLICANT: Hook, Magnus
; APPLICANT: Patti, Joseph M.
; APPLICANT: House-Pompeo, Karen
; APPLICANT: Sthanam, Narayana
; APPLICANT: Symersky, Jindrich
; TITLE OF INVENTION: COLLAGEN BINDING PROTEIN COMPOSITIONS
; TITLE OF INVENTION: AND METHODS OF USE
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Arnold, White & Durkee
; STREET: P.O. Box 4433
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.
; ZIP: 77210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/856,253
; FILING DATE: Concurrently Herewith
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/017,678
; FILING DATE: 16-MAY-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Kitchell, Barbara S.
; REGISTRATION NUMBER: 33,928
; REFERENCE/DOCKET NUMBER: TAMK:193
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (512) 418-3000
; TELEFAX: (512) 474-7577
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 211 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
US-08-856-253-4

Query Match 90.5%; Score 19; DB 4; Length 211;
Best Local Similarity 66.7%; Pred. No. 4.le+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
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Db 34 EAGTSS 39

RESULT 4
US-08-405-175A-5
; Sequence 5, Application US/08405175A
; Patent No. 5885772
; GENERAL INFORMATION:
; APPLICANT: Aderem, Alan A.
; APPLICANT: Chen, Jianmin
; APPLICANT: Chang, Sandy
; TITLE OF INVENTION: METHOD FOR THE DETECTION OF ANENCEPHALY
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Klauber & Jackson
; STREET: 411 Hackensack Avenue

;; CITY: Hackensack
;; STATE: New Jersey
;; COUNTRY: USA
;; ZIP: 07601
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;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.25
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/405,175A
;; FILING DATE: 16-MAR-1995
;; CLASSIFICATION: 435
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Jackson Esq., David A.
;; REGISTRATION NUMBER: 26,742
;; REFERENCE/DOCKET NUMBER: 600-1-121A
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 201 487-5800
;; TELEFAX: 201 343-1684
;; TELEX: 133521
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;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 332 amino acids
;; TYPE: amino acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: protein
;; DESCRIPTION: predicted primary structure of human MARCKS
;; HYPOTHETICAL: NO
;; US-08-405-175A-5

Query Match 90.5%; Score 19; DB 2; Length 332;
Best Local Similarity 66.7%; Pred. No. 6.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
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Db 206 EAGAS 211

RESULT 5
US-08-191-866D-21
; Sequence 21, Application US/08191866D
; Patent No. 5783195
; GENERAL INFORMATION:
; APPLICANT: Cochran, Mark D.
; TITLE OF INVENTION: Recombinant Infectious Bovine
; TITLE OF INVENTION: Rhinotracheitis Virus S-IBR-052 And Uses Thereof
; NUMBER OF SEQUENCES: 99
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John P. White
; STREET: 1185 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/191,866D
; FILING DATE: 4 February 1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: White, John P.
; REGISTRATION NUMBER: 28,678
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 278-0400

;; TELEFAX: (212) 391-0525
;; TELEX: 422523
;; INFORMATION FOR SEQ ID NO: 21:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 441 amino acids
;; TYPE: amino acid
;; TOPOLOGY: linear
;; MOLECULE TYPE: protein
;; US-08-191-866D-21

Query Match 90.5%; Score 19; DB 1; Length 441;
Best Local Similarity 66.7%; Pred. No. 8.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
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Db 265 EAGSAS 270

RESULT 6
US-08-185-949B-21
; Sequence 21, Application US/08185949B
; Patent No. 5874279
; GENERAL INFORMATION:
; APPLICANT: Mark D. Cochran
; APPLICANT: Richard D. Macdonald
; TITLE OF INVENTION: Recombinant Infectious Bovine
; TITLE OF INVENTION: Rhinotracheitis Virus
; NUMBER OF SEQUENCES: 104
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John P. White
; STREET: 1185 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM 330 466 DX2
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/185,949B
; FILING DATE: 03-NOV-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: White, John P.
; REGISTRATION NUMBER: 678
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 278-0400
; TELEFAX: (212) 278-0525
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 441 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-185-949B-21

Query Match 90.5%; Score 19; DB 2; Length 441;
Best Local Similarity 66.7%; Pred. No. 8.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
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Db 265 EAGSAS 270

RESULT 7
US-08-856-253-6
; Sequence 6, Application US/08856253

; Patent No. 6288214
; GENERAL INFORMATION:
; APPLICANT: Hook, Magnus
; APPLICANT: Patti, Joseph M.
; APPLICANT: House-Pompeo, Karen
; APPLICANT: Sthanam, Narayana
; APPLICANT: Symersky, Jindrich
; TITLE OF INVENTION: COLLAGEN BINDING PROTEIN COMPOSITIONS
; TITLE OF INVENTION: AND METHODS OF USE
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Arnold, White & Durkee
; STREET: P.O. Box 4433
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.
; ZIP: 77210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/856.253
; FILING DATE: Concurrently Herewith
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/017,678
; FILING DATE: 16-MAY-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Kitchell, Barbara S.
; REGISTRATION NUMBER: 33,928
; REFERENCE/DOCKET NUMBER: TAMK.193
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (512) 418-3000
; TELEFAX: (512) 474-7577
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 512 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; US-08-856-253-6

Query Match 90.5%; Score 19; DB 4; Length 512;
Best Local Similarity 66.7%; Pred. No. 1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
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Db 149 EAGTSS 154

RESULT 8
PCT-US95-03747-3
; Sequence 3, Application PC/TUS9503747
; GENERAL INFORMATION:
; APPLICANT: LA JOLLA CANCER RESEARCH FOUNDATION
; TITLE OF INVENTION: Brevican, A Glial Cell Proteoglycan
; NUMBER OF SEQUENCES: 3
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/03747
; FILING DATE: 27-MAR-1995
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Imbra, Richard J.
; REGISTRATION NUMBER: 37,643
; REFERENCE/DOCKET NUMBER: FP-LJ 1453
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 908 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; PCT-US95-03747-3

Query Match 90.5%; Score 19; DB 5; Length 908;
Best Local Similarity 66.7%; Pred. No. 1.8e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
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Db 583 EAGSSS 588

RESULT 9
US-09-130-242-2
; Sequence 2, Application US/09130242B
; Patent No. 6194558
; GENERAL INFORMATION:
; APPLICANT: Gianturco, S.H.
; APPLICANT: Bradley, W.A.
; TITLE OF INVENTION: DNA Encoding Human Monocyte-Macrophage Aoplipoprotein
; FILE REFERENCE: D5880
; CURRENT APPLICATION NUMBER: US/09/130,242B
; CURRENT FILING DATE: 1998-08-06
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: MS WORD, Macintosh OS 8.5
; SEQ ID NO 2
; LENGTH: 1088
; TYPE: PRT
; ORGANISM: Homo sapien
; US-09-130-242-2

Query Match 90.5%; Score 19; DB 4; Length 1088;
Best Local Similarity 66.7%; Pred. No. 2.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
| | | | |
Db 327 EAGTAS 332

RESULT 10
US-08-447-031A-2
; Sequence 2, Application US/08447031A
; Patent No. 5851794
; GENERAL INFORMATION:
; APPLICANT: GUSS, Bengt
; APPLICANT: HOOK, Magnus
; APPLICANT: JONSSON, Hans
; APPLICANT: LINDBERG, Martin
; APPLICANT: PATTI, Joseph
; APPLICANT: SIGNAS, Christer
; TITLE OF INVENTION: A COLLAGEN BINDING PROTEIN AS WELL AS
; TITLE OF INVENTION: ITS PREPARATION
; NUMBER OF SEQUENCES: 8

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Burns, Doane, Swecker & Mathis
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/447,031A
; FILING DATE: 22-MAY-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/861,804
; FILING DATE: 21-AUG-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/SE91/00707
; FILING DATE: 22-OCT-1991
; APPLICATION DATA: SE 9003374-7
; FILING DATE: 22-OCT-1990
; ATTORNEY/AGENT INFORMATION:
; NAME: McGowan, Malcolm K.
; REGISTRATION NUMBER: 39,300
; REFERENCE/DOCKET NUMBER: 012889-006
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1183 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-447-031A-2

Query Match 90.5%; Score 19; DB 2; Length 1183;
Best Local Similarity 66.7%; Pred. No. 2.3e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 166 EAGTSS 171

RESULT 11
US-08-485-355B-31
; Sequence 31, Application US/08485355B
; Patent No. 6177075
; GENERAL INFORMATION:
; APPLICANT: Christian, P. D., Gordon, K. H.J., Hanzlik, T. N.
; TITLE OF INVENTION: Insect Viruses and Their Uses in
; Protecting Plants
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Flehr Hobbach Test Albritton & Herbert LLP
; STREET: Four Embarcadero Center, Suite 3400
; CITY: San Francisco
; STATE: California
; COUNTRY: United States
; ZIP: 94111-4187
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/485,355B

; FILING DATE: 07-Jun-1995
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/440,522
; FILING DATE: 12-MAY-1995
; APPLICATION NUMBER: US 08/089,372
; FILING DATE: 08-JUL-1993
; APPLICATION NUMBER: AU PL4081/92
; FILING DATE: 14-AUG-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Irecartin, Richard F.
; REGISTRATION NUMBER: 31,801
; REFERENCE/DOCKET NUMBER: A-58631-2/RFT/DSS
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 781-1989
; TELEFAX: (415) 398-3249
; TELEX: 910 277299
; INFORMATION FOR SEQ ID NO: 31:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; SEQUENCE DESCRIPTION: SEQ ID NO: 31:
US-08-485-355B-31

Query Match 85.7%; Score 18; DB 4; Length 9;
Best Local Similarity 66.7%; Pred. No. 1.7e+05;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 3 EAGVAS 8

RESULT 12
US-08-596-387B-109
; Sequence 109, Application US/08596387B
; Patent No. 5869270
; GENERAL INFORMATION:
; APPLICANT: Rhode, Peter R.
; APPLICANT: Jiao, Jin-An
; APPLICANT: Burkhardt, Martin
; APPLICANT: Wong, Hing
; TITLE OF INVENTION: MHC COMPLEXES AND USES THEREOF
; NUMBER OF SEQUENCES: 124
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Dade International, Inc.
; STREET: 1717 Deerfield Road
; CITY: Deerfield
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60015
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/596,387B
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/09816
; FILING DATE: 31-JUL-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/382,454
; FILING DATE: 01-FEB-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/283,302
; FILING DATE: 29-JUL-1994
; ATTORNEY/AGENT INFORMATION:

NAME: Pearson, Louise S.
REGISTRATION NUMBER: 32,369
REFERENCE/DOCKET NUMBER: STR-4665-CIP2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (708) 267-5300
TELEFAX: (708) 267-5376
INFORMATION FOR SEQ ID NO: 109:
SEQUENCE CHARACTERISTICS:
LENGTH: 59 amino acids
TYPE: amino acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
US-08-596-387B-109

Query Match 85.7%; Score 18; DB 2; Length 59;
Best Local Similarity 66.7%; Pred. No. 2.2e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 41 EAGRAS 46

RESULT 13
US-09-067-615-109
; Sequence 109, Application US/09067615
; Patent No. 6309645
; GENERAL INFORMATION:
; APPLICANT: Rhode, Peter R.
; APPLICANT: Jiao, Jin-An
; APPLICANT: Burkhardt, Martin
; APPLICANT: Wong, Hing
; TITLE OF INVENTION: MHC COMPLEXES AND USES THEREOF
; NUMBER OF SEQUENCES: 124
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dade International, Inc.
; STREET: 1717 Deerfield Road
; CITY: Deerfield
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60015
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/067,615
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/596,387
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/382,454
; FILING DATE: 01-FEB-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/283,302
; FILING DATE: 29-JUL-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Pearson, Louise S.
; REGISTRATION NUMBER: 32,369
; REFERENCE/DOCKET NUMBER: STR-4665-CIP2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (708) 267-5300
; TELEFAX: (708) 267-5376
; INFORMATION FOR SEQ ID NO: 109:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 59 amino acids
; TYPE: amino acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown

US-09-067-615-109

Query Match 85.7%; Score 18; DB 4; Length 59;
Best Local Similarity 66.7%; Pred. No. 2.2e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 41 EAGRAS 46

RESULT 14
PCT-US95-09816A-109
; Sequence 109, Application PC/TUS9509816A
; GENERAL INFORMATION:
; APPLICANT: Wong, Hing C.
; APPLICANT: Rhode, Peter R.
; APPLICANT: Widanz, Jon A.
; APPLICANT: Grammer, Susan
; APPLICANT: Edwards, Ana C.
; APPLICANT: Chavaillaz, Pierre-Andre
; APPLICANT: Jiao, Jin-An
; TITLE OF INVENTION: MHC COMPLEXES AND USES THEREOF
; NUMBER OF SEQUENCES: 123
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dade International, Inc.
; STREET: 1717 Deerfield Road
; CITY: Deerfield
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60015
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/09816A
; FILING DATE: 31-JUL-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/382,454
; FILING DATE: 01-FEB-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/283,302
; FILING DATE: 29-JUL-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Pearson, Louise S.
; REGISTRATION NUMBER: 32,369
; REFERENCE/DOCKET NUMBER: STR-4665-CIP2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (708) 267-5300
; TELEFAX: (708) 267-5376
; INFORMATION FOR SEQ ID NO: 109:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 59 amino acids
; TYPE: amino acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; PCT-US95-09816A-109

Query Match 85.7%; Score 18; DB 5; Length 59;
Best Local Similarity 66.7%; Pred. No. 2.2e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 41 EAGRAS 46

RESULT 15

US-08-446-137B-12
; Sequence 12, Application US/08446137B
; Patent No. 6162903
; GENERAL INFORMATION:
; APPLICANT: Trowern, Angus R.
; APPLICANT: Atkinson, Anthony
; APPLICANT: Murphy, Jonathan P.
; APPLICANT: Laurence, Oliver S.
; APPLICANT: Duggleby, Clive J.
; TITLE OF INVENTION: IMMUNOGLOBULIN BINDING PROTEINS DERIVED
; FROM L PROTEIN AND THEIR USES
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED AND BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/446,137B
; FILING DATE: 22-MAY-1995
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Mcmasters, David D.
; REGISTRATION NUMBER: 33,963
; REFERENCE/DOCKET NUMBER: 100084.406
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 61 amino acids
; TYPE: amino acid
; STRANDEDNESS: linear
; TOPOLOGY: linear
US-08-446-137B-12

Query Match 85.7%; Score 18; DB 4; Length 61;
Best Local Similarity 66.7%; Pred. No. 2.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 29 EAGTGS 34

RESULT 16
US-08-446-137B-10
; Sequence 10, Application US/08446137B
; Patent No. 6162903
; GENERAL INFORMATION:
; APPLICANT: Trowern, Angus R.
; APPLICANT: Atkinson, Anthony
; APPLICANT: Murphy, Jonathan P.
; APPLICANT: Laurence, Oliver S.
; APPLICANT: Duggleby, Clive J.
; TITLE OF INVENTION: IMMUNOGLOBULIN BINDING PROTEINS DERIVED
; FROM L PROTEIN AND THEIR USES
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED AND BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/446,137B
; FILING DATE: 22-MAY-1995
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Mcmasters, David D.
; REGISTRATION NUMBER: 33,963
; REFERENCE/DOCKET NUMBER: 100084.406
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 66 amino acids
; TYPE: amino acid
; STRANDEDNESS: linear
; TOPOLOGY: linear
US-08-446-137B-10

Query Match 85.7%; Score 18; DB 4; Length 66;
Best Local Similarity 66.7%; Pred. No. 2.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 33 EAGTGS 38

RESULT 17
US-08-598-873-19
; Sequence 19, Application US/08598873
; Patent No. 5928884
; GENERAL INFORMATION:
; APPLICANT: Croce, Carlo M.
; APPLICANT: Huebner, Kay
; TITLE OF INVENTION: PHIT PROTEINS AND NUCLEIC ACIDS AND
; METHODS BASED THEREON
; NUMBER OF SEQUENCES: 77
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: U.S.A.
; ZIP: 10036-2711
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/598,873
; FILING DATE: 09-FEB-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Friebe, Thomas E.
; REGISTRATION NUMBER: 29,258
; REFERENCE/DOCKET NUMBER: 8666-004
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 790-9090
; TELEFAX: (212) 869-9741/8864
; TELETYPE: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 91 amino acids
; TYPE: amino acid
; TOPOLOGY: unknown

; MOLECULE TYPE: peptide
US-08-598-873-19

Query Match 85.7%; Score 18; DB 2; Length 91;
Best Local Similarity 66.7%; Pred. No. 3.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 49 EAGKSS 54

RESULT 18

US-08-605-430-19
Sequence 19, Application US/08605430
Patent No. 6242212

GENERAL INFORMATION:

APPLICANT: Croce, Carlo M.
APPLICANT: Huebner, Kay
TITLE OF INVENTION: PHIT PROTEINS AND NUCLEIC ACIDS AND
METHODS BASED THEREON
NUMBER OF SEQUENCES: 86
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: U.S.A.
ZIP: 10036-2711

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA: US/08/605,430

APPLICATION NUMBER: US/08/605,430

FILING DATE: 22-FEB-1996

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: Friebe, Thomas E.
REGISTRATION NUMBER: 29,258
REFERENCE/DOCKET NUMBER: 8666-005
TELEPHONE: (212) 790-9090
TELEFAX: (212) 869-9741/8864
TELEX: 66141 PENNIE

INFORMATION FOR SEQ ID NO: 19:

SEQUENCE CHARACTERISTICS:

LENGTH: 91 amino acids
TYPE: amino acid
TOPOLOGY: unknown
MOLECULE TYPE: peptide
US-08-605-430-19

Query Match 85.7%; Score 18; DB 4; Length 91;
Best Local Similarity 66.7%; Pred. No. 3.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 49 EAGKSS 54

RESULT 19

US-07-869-912-2

Sequence 2, Application US/07869912
Patent No. 5316922

GENERAL INFORMATION:

APPLICANT: Court, Don
APPLICANT: Brown, Stanley
TITLE OF INVENTION: A Method for Identifying and

; TITLE OF INVENTION: Expressing Proteins that Recognize and Adhere to Specific
; TITLE OF INVENTION: Probes
; NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Kenneth A. Weber
STREET: One Market Plaza, Steuart Tower, Suite 2000
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94105

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/869,912

FILING DATE:

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Weber, Kenneth A.
REGISTRATION NUMBER: 32,334
REFERENCE/DOCKET NUMBER: 15280-10
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-543-9600
TELEFAX: 415-543-5043
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 123 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-07-869-912-2

Query Match 85.7%; Score 18; DB 1; Length 123;
Best Local Similarity 66.7%; Pred. No. 4.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 21 EAGGSS 26

RESULT 20

US-08-446-137B-9

Sequence 9, Application US/08446137B

Patent No. 6162903

GENERAL INFORMATION:

APPLICANT: Trower, Angus R.
APPLICANT: Atkinson, Anthony
APPLICANT: Murphy, Jonathan P.
APPLICANT: Laurence, Oliver S.
APPLICANT: Dugleby, Clive J.

TITLE OF INVENTION: IMMUNOGLOBULIN BINDING PROTEINS DERIVED

FROM L PROTEIN AND THEIR USES

NUMBER OF SEQUENCES: 12

CORRESPONDENCE ADDRESS:

ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/446,137B
FILING DATE: 22-MAY-1995
CLASSIFICATION: 514

ATTORNEY/AGENT INFORMATION:
NAME: McMasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 100084.406
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 175 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-446-137B-9

Query Match 85.7%; Score 18; DB 4; Length 175;
Best Local Similarity 66.7%; Pred. No. 6.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
||| |
Db 29 EAGITS 34

RESULT 21
US-08-708-958-2
Sequence 2, Application US/08708958
Patent No. 5948952

GENERAL INFORMATION:
APPLICANT: SANDS, Arthur T.
APPLICANT: BRADLEY, Allan
APPLICANT: ABUIN, Alejandro
TITLE OF INVENTION: XERODERMA PIGMENTOSUM-DEFICIENT
TITLE OF INVENTION: MOUSE
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS
STREET: 2100 PENNSYLVANIA AVENUE, N.W.
CITY: WASHINGTON
STATE: D.C.
COUNTRY: USA
ZIP: 20037

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
FILING DATE: SEP-1996
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: KIT, Gordon
REGISTRATION NUMBER: 30,764
REFERENCE/DOCKET NUMBER: A-6641
TELEPHONE: (202) 293-7060
TELEFAX: (202) 293-7860
TELEX: 6491103

INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 211 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-708-958-2

Query Match 85.7%; Score 18; DB 2; Length 211;
Best Local Similarity 66.7%; Pred. No. 7.8e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
||| |
Db 25 EAGSGS 30

RESULT 22
US-09-423-340-2
Sequence 2, Application US/09423340
Patent No. 6225454
GENERAL INFORMATION:
APPLICANT: MIYAGI, Taeko
APPLICANT: WADA, Tadashi
APPLICANT: YOSHIKAWA, Yuko
TITLE OF INVENTION: SIALIDASE LOCALIZED IN PLASMA MEMBRANE AND
FILE REFERENCE: OP699
CURRENT APPLICATION NUMBER: US/09/423,340
CURRENT FILING DATE: 1999-11-22
EARLIER APPLICATION NUMBER: JP 9-132174
EARLIER FILING DATE: 1997-05-22
NUMBER OF SEQ ID NOS: 24
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 2
LENGTH: 428
TYPE: PRT
ORGANISM: Bos primigenius
US-09-423-340-2

Query Match 85.7%; Score 18; DB 4; Length 428;
Best Local Similarity 66.7%; Pred. No. 1.6e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
||| |
Db 321 EAGTSL 326

RESULT 23
US-08-749-902-1
Sequence 1, Application US/08749902
Patent No. 5985635
GENERAL INFORMATION:
APPLICANT: Bandman, Olga
APPLICANT: Goli, Surva K.
APPLICANT: Hillman, Jennifer L.
TITLE OF INVENTION: NOVEL HUMAN SERINE/THREONINE
TITLE OF INVENTION: PROTEIN KINASES
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: US
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/749,902
FILING DATE: Filed Herewith
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0150 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555

TELEFAX: 415-845-4166
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 433 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
CLONE: Consensus
US-08-749-902-1

Query Match 85.7%; Score 18; DB 2; Length 433;
Best Local Similarity 66.7%; Pred. No. 1.6e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eaqxss 6
||| |
Db 396 EAGAVS 401

RESULT 24
US-09-330-095-1
Sequence 1, Application US/09330095
Patent No. 6127161
GENERAL INFORMATION:
APPLICANT: Kikkoman Corporation
TITLE OF INVENTION: Leucine Aminopeptidase Gene, Recombinant DNA, and
FILE REFERENCE: PH-622
CURRENT APPLICATION NUMBER: US/09/330,095
CURRENT FILING DATE: 1999-06-11
EARLIER APPLICATION NUMBER: JP-164611/1998
EARLIER FILING DATE: 1998-06-12
NUMBER OF SEQ ID NOS: 14
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 1
LENGTH: 481
TYPE: PRT
ORGANISM: Aspergillus sojae
US-09-330-095-1

Query Match 85.7%; Score 18; DB 3; Length 481;
Best Local Similarity 66.7%; Pred. No. 1.8e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eaqxss 6
||| |
Db 193 EAGSVS 198

RESULT 25
US-08-960-190A-25
Sequence 25, Application US/08960190A
Patent No. 6232445
GENERAL INFORMATION:
APPLICANT: Rhode, Peter R.
APPLICANT: Acevedo, Jorge
APPLICANT: Burkhardt, Martin
APPLICANT: Jiao, Jin-an
APPLICANT: Wong, Hing C.
TITLE OF INVENTION: SOLUBLE MHC COMPLEXES AND
METHODS OF USE THEREOF
NUMBER OF SEQUENCES: 38
CORRESPONDENCE ADDRESS:
ADDRESSEE: Dike, Bronstein, Roberts & Cushman, LLP
STREET: 130 Water Street
CITY: Boston
STATE: MA
COUNTRY: usa
ZIP: 02109
COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/960,190A
FILING DATE: 29-OCT-1997
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Corless, Peter F
REGISTRATION NUMBER: 33,860
REFERENCE/DOCKET NUMBER: 48002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-523-3400
TELEFAX: 617-523-6440
TELEX:

INFORMATION FOR SEQ ID NO: 25:
SEQUENCE CHARACTERISTICS:
LENGTH: 500 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
FRAGMENT TYPE: internal
US-08-960-190A-25

Query Match 85.7%; Score 18; DB 4; Length 500;
Best Local Similarity 66.7%; Pred. No. 1.8e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eaqxss 6
||| |
Db 41 EAGRAS 46

RESULT 26
US-07-612-673-2
Sequence 2, Application US/07612673
Patent No. 5260208
GENERAL INFORMATION:
APPLICANT: Petre, Dominique
APPLICANT: Cerbelaud, Edith
APPLICANT: Mayaux, Jean-Francois
APPLICANT: Yeh, Patrice
TITLE OF INVENTION: NOVEL POLYPEPTIDES, THE DNA SEQUENCES
TITLE OF INVENTION: ALLOWING THEIR EXPRESSION, METHOD OF PREPARATION, AND
NUMBER OF SEQUENCES: 17
CORRESPONDENCE ADDRESS:
ADDRESSEE: Finnegan, Henderson, Farabow, Garrett &
ADDRESSEE: Dunner
STREET: 1300 I Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20005
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/612,673
FILING DATE: 19901114
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Potter, Jane E.R.
REGISTRATION NUMBER: 33,332
REFERENCE/DOCKET NUMBER: 03715.0010
TELECOMMUNICATION INFORMATION:

TELEPHONE: (202) 408-4000
TELEFAX: (202) 408-4400
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 503 amino acids
TYPE: AMINO ACID
TOPOLOGY: linear
MOLECULE TYPE: protein
US-07-612-673-2

Query Match 85.7%; Score 18; DB 1; Length 503;
Best Local Similarity 66.7%; Pred. No. 1.8e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
||| |

Db 149 EAGSS 154

RESULT 27
US-08-845-258-52
Sequence 52, Application US/08845258
Patent No. 6183976
GENERAL INFORMATION:

APPLICANT: Reed, Steven G.
APPLICANT: Lodes, Michael J.
APPLICANT: Houghton, Raymond
APPLICANT: Sleath, Paul R.
TITLE OF INVENTION: COMPOUNDS AND METHODS FOR THE DIAGNOSIS
AND TREATMENT OF B. MICROTI INFECTION
NUMBER OF SEQUENCES: 53
CORRESPONDENCE ADDRESS:

ADDRESSEE: SEED AND BERRY
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/845,258
FILING DATE: 24-APR-1997

CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Maki, David J.
REGISTRATION NUMBER: 31,392
REFERENCE/DOCKET NUMBER: 210121.426C1
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031

INFORMATION FOR SEQ ID NO: 52:
SEQUENCE CHARACTERISTICS:
LENGTH: 503 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Babesia Microti
US-08-845-258-52

Query Match 85.7%; Score 18; DB 4; Length 503;
Best Local Similarity 66.7%; Pred. No. 1.8e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
||| |

Db 417 EAGTS 422

RESULT 28
US-08-990-571-52
Sequence 52, Application US/08990571
Patent No. 6214971
GENERAL INFORMATION:

APPLICANT: Reed, Steven G. et al.
TITLE OF INVENTION: COMPOUNDS AND METHODS FOR THE DIAGNOSIS AND TREATMENT OF B.
NUMBER OF SEQUENCES: 79
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED AND BERRY
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/990,571
FILING DATE: 11-DEC-1997

CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Maki, David J.
REGISTRATION NUMBER: 31,392
REFERENCE/DOCKET NUMBER: 210121.426C2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031

INFORMATION FOR SEQ ID NO: 52:
SEQUENCE CHARACTERISTICS:
LENGTH: 503 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Babesia Microti
US-08-990-571-52

Query Match 85.7%; Score 18; DB 4; Length 503;
Best Local Similarity 66.7%; Pred. No. 1.8e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
||| |

Db 417 EAGTS 422

RESULT 29
US-07-796-361A-11
Sequence 11, Application US/07796361A
Patent No. 5258292
GENERAL INFORMATION:

APPLICANT: YEH, Patrice
APPLICANT: MAYAUX, Jean-Francois
APPLICANT: CERBELAUD, Edith
APPLICANT: PETRE, Dominique
TITLE OF INVENTION: ENZYMIC PROCESS FOR THE SYNTHESIS OF
NUMBER OF SEQUENCES: 24
CORRESPONDENCE ADDRESS:
ADDRESSEE: Burns, Doane, Swecker and Mathis
STREET: The George Mason Building, Washington &
CITY: Prince Streets
STATE: Alexandria
CITY: Alexandria
STATE: Virginia

; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/796.361A
; FILING DATE: 19911122
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 90-14 853
; FILING DATE: 28-NOV-1990
; ATTORNEY/AGENT INFORMATION:
; NAME: CRANE-FEURY, SHARON E.
; REGISTRATION NUMBER: P36,113
; REFERENCE/DOCKET NUMBER: 003025-010
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; TELEX: 440580
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 521 amino acids
; TYPE: AMINO ACID
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-07-796-361A-11

Query Match 85.7%; Score 18; DB 1; Length 521;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 eagxxs 6
||| |
Db 167 EAGGS 172

RESULT 30
US-08-539-666-2
; Sequence 2, Application US/08539666
; Patent No. 5766918
; GENERAL INFORMATION:
; APPLICANT: Petre, Dominique
; APPLICANT: Cerbelaud, Edith
; APPLICANT: Mayaux, Jean-Francois
; APPLICANT: Yeh, Patrice
; TITLE OF INVENTION: No. 5766918el Polypeptides, The DNA Sequences
; TITLE OF INVENTION: Allowing their Expression, Method of Preparation, and
; TITLE OF INVENTION: Utilization
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Finnegan, Henderson, Farabow, Garrett &
; ADDRESSEE: Dunner
; STREET: 1300 I Street, N.W.
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3315
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/539,666
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/097,009
; FILING DATE: 27-JUL-1993

; APPLICATION NUMBER: US 07/612,673
; FILING DATE: 14-NOV-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 8916332
; FILING DATE: 11-DEC-1989
; ATTORNEY/AGENT INFORMATION:
; NAME: Potter, Jane E.R.
; REGISTRATION NUMBER: 33,332
; REFERENCE/DOCKET NUMBER: 03715.0010-01000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-408-4000
; TELEFAX: 202-408-4400
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 521 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-539-666-2

Query Match 85.7%; Score 18; DB 1; Length 521;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 1 eagxxs 6
||| |
Db 167 EAGGS 172

RESULT 31
US-08-348-891A-2
; Sequence 2, Application US/08348891A
; Patent No. 5654136
; GENERAL INFORMATION:
; APPLICANT: SASAKI, Keiko
; APPLICANT: MORI, Takayuki
; APPLICANT: MAKINO, Satoshi
; TITLE OF INVENTION: ATTENUATED MEASLES VIRUS VACCINE,
; TITLE OF INVENTION: CONTAINING SPECIFIC NUCLEOTIDE SEQUENCE AND A METHOD FOR
; TITLE OF INVENTION: ITS ABSOLUTE IDENTIFICATION
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: YOUNG & THOMPSON
; STREET: 745 South 23rd Street
; CITY: Arlington
; STATE: Virginia
; COUNTRY: USA
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/348,891A
; FILING DATE: 25-NOV-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/848,400
; FILING DATE: 10-MAR-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: JP 3-293625
; FILING DATE: 14-OCT-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: PATCH, Andrew J.
; REGISTRATION NUMBER: 32,925
; REFERENCE/DOCKET NUMBER: KP-7501
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 703-521-2297
; TELEFAX: 703-685-0573
; TELEX: 248425 EMBON
; INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:
LENGTH: 525 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-348-891A-2

Query Match 85.7%; Score 18; DB 1; Length 525;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxvs 6
||| |
DB 263 EAGLAS 268

RESULT 32

US-08-905-817-2
Sequence 2, Application US/08905817
Patent No. 5824777

GENERAL INFORMATION:
APPLICANT: SASAKI, Keiko
APPLICANT: MAKINO, Satoshi
TITLE OF INVENTION: ATTENUATED MEASLES VIRUS VACCINE,
TITLE OF INVENTION: CONTAINING SPECIFIC NUCLEOTIDE SEQUENCE AND A METHOD FOR
TITLE OF INVENTION: ITS ABSOLUTE IDENTIFICATION
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: YOUNG & THOMPSON
STREET: 745 South 23rd Street
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22202

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/905,817
FILING DATE: 04-AUG-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/348,891
FILING DATE: 25-NOV-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/848,400
FILING DATE: 10-MAR-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: JP 3-293625
FILING DATE: 14-OCT-1991
ATTORNEY/AGENT INFORMATION:
NAME: PATCH, Andrew J.
REGISTRATION NUMBER: 32,925
REFERENCE/DOCKET NUMBER: KP-7501A
TELECOMMUNICATION INFORMATION:
TELEPHONE: 703-521-2297
TELEFAX: 703-685-0573
TELEX: 248425 EMBON
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 525 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-905-817-2

Query Match 85.7%; Score 18; DB 2; Length 525;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 eagxvs 6
||| |
DB 263 EAGLAS 268

RESULT 33

US-08-513-841-1
Sequence 1, Application US/08513841
Patent No. 5753481

GENERAL INFORMATION:
APPLICANT: Niwa, Mineo
APPLICANT: Saito, Yoshimasa
APPLICANT: Ishii, Yoshinori
APPLICANT: Yoshida, Masaru
APPLICANT: Suzuki, Hiroshi
TITLE OF INVENTION: No. 5753481el L-sorbose Dehydrogenase and No. 5753481el L-s
TITLE OF INVENTION: Dehydrogenase Obtained from Gluconobacter oxydans T-100
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Oblon, Spivak, McClelland, Maier & Neustadt, P.C.
STREET: 1755 Jefferson Davis Highway, Suite 400
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22202

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44 Mb storage
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: MS-DOS Editor
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/513,841
FILING DATE: 01-NOV-1995
CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: UK 9304700.9
FILING DATE: 08-MAR-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: JP 241851/1993
FILING DATE: 28-SEP-1993
ATTORNEY/AGENT INFORMATION:
NAME: NORMAN F. OBLON
REGISTRATION NUMBER: 24,618
REFERENCE/DOCKET NUMBER: 18-909-0 PCT

TELECOMMUNICATION INFORMATION:
TELEPHONE: 703-413-3000
TELEFAX: 703-413-2220
TELEX: 248855 OPAT UR
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 530 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
ORIGINAL SOURCE:
ORGANISM: Gluconobacter oxydans
STRAIN: T-100
FEATURE:
NAME/KEY: mat peptide
LOCATION: 1..530
IDENTIFICATION METHOD: experimentally
US-08-513-841-1

Query Match 85.7%; Score 18; DB 1; Length 530;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxvs 6
||| |
DB 364 EAGVTS 369

RESULT 34
US-08-696-834-1
; Sequence 1, Application US/08696834
; Patent No. 5834263
; GENERAL INFORMATION:
; APPLICANT: Niwa, Mineo
; APPLICANT: Saito, Yoshimasa
; APPLICANT: Ishii, Yoshinori
; APPLICANT: Yoshida, Masaru
; APPLICANT: Hayashi, Hiromi
; TITLE OF INVENTION: Method for Producing 2-Keto-L-Gulonic Acid
; NUMBER OF SEQUENCES: 48
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Obion, Spivak, McClelland, Maier & Neustadt,
; STREET: 1755 Jefferson Davis Highway, Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: USA
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette - 3.50 inch, 1.44 Mb storage
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE:
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/696,834
; FILING DATE: 24-SEP-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: JP 28612/1994
; FILING DATE: 25-FEB-1994
; ATTORNEY/AGENT INFORMATION:
; NAME:
; REGISTRATION NUMBER:
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 413-3000
; TELEFAX: (703) 413-2220
; TELEX: 248855 OPAT UR
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 530 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; ORIGINAL SOURCE:
; ORGANISM: Gluconobacter oxydans
; STRAIN: T-100
; FEATURE:
; NAME/KEY: mat peptide
; LOCATION: 1..530
; IDENTIFICATION METHOD: experimentally
; US-08-696-834-1

Query Match 85.7%; Score 18; DB 2; Length 530;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
Db 364 EAGVTS 369

RESULT 35
US-08-942-673-1
; Sequence 1, Application US/08942673
; Patent No. 5861292
; GENERAL INFORMATION:
; APPLICANT: Niwa, Mineo

; APPLICANT: Saito, Yoshimasa
; APPLICANT: Ishii, Yoshinori
; APPLICANT: Yoshida, Masaru
; APPLICANT: Suzuki, Hiromi
; TITLE OF INVENTION: No. 5861292el L-sorbose Dehydrogenase and No. 5861292el
; TITLE OF INVENTION: L-sorbose Dehydrogenase Obtained from Gluconobacter
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Obion, Spivak, McClelland, Maier & Neustadt, P.C.
; STREET: 1755 Jefferson Davis Highway, Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: USA
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.44 Mb storage
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: MS-DOS Editor
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/942,673
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/513,841
; FILING DATE: 01-NOV-1995
; APPLICATION NUMBER: UK 9304700.9
; FILING DATE: 08-MAR-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: JP 241851/1993
; FILING DATE: 28-SEP-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: NORMAN F. OBLON
; REGISTRATION NUMBER: 24,618
; REFERENCE/DOCKET NUMBER: 18-909-0 PCT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 703-413-3000
; TELEFAX: 703-413-2220
; TELEX: 248855 OPAT UR
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 530 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; ORIGINAL SOURCE:
; ORGANISM: Gluconobacter oxydans
; STRAIN: T-100
; FEATURE:
; NAME/KEY: mat peptide
; LOCATION: 1..530
; IDENTIFICATION METHOD: experimentally
; US-08-942-673-1

Query Match 85.7%; Score 18; DB 2; Length 530;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
Db 364 EAGVTS 369

RESULT 36
US-09-118-317-1
; Sequence 1, Application US/09118317
; Patent No. 6197562
; GENERAL INFORMATION:
; APPLICANT: Niwa, Mineo
; APPLICANT: Saito, Yoshimasa
; APPLICANT: Ishii, Yoshinori

APPLICANT: Yoshida, Masaru
APPLICANT: Suzuki, Hiromi
TITLE OF INVENTION: No. 6197562el L-sorbose Dehydrogenase and No. 6197562el
TITLE OF INVENTION: L-sorbose Dehydrogenase Obtained from Gluconobacter
TITLE OF INVENTION: oxydans T-100
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Oblon, Spivak, McClelland, Maier & Neustadt, P.C.
STREET: 1755 Jefferson Davis Highway, Suite 400
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44 Mb storage
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: MS-DOS Editor
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/118,317
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/513,841
FILING DATE: 01-NOV-1995
APPLICATION NUMBER: UK 9304700.9
FILING DATE: 08-MAR-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: JP 241851/1993
FILING DATE: 28-SEP-1993
ATTORNEY/AGENT INFORMATION:
NAME: NORMAN F. OBLON
REGISTRATION NUMBER: 24,618
REFERENCE/DOCKET NUMBER: 18-909-0 PCT
TELECOMMUNICATION INFORMATION:
TELEPHONE: 703-413-3000
TELEFAX: 703-413-2220
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 530 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
ORIGINAL SOURCE:
ORGANISM: Gluconobacter oxydans
STRAIN: T-100
FEATURE:
NAME/KEY: mat peptide
LOCATION: 1.530
IDENTIFICATION METHOD: experimentally
US-09-118-317-1

Query Match 85.7%; Score 18; DB 4; Length 530;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
Db 364 EAGVTS 369

RESULT 37
US-08-808-931-18
Sequence 18, Application US/08808931
Patent No. 5939602
GENERAL INFORMATION:
APPLICANT: Volrath, Sandra
APPLICANT: Johnson, Marie
APPLICANT: Potter, Sharon
APPLICANT: Ward, Eric
APPLICANT: Heifetz, Peter

TITLE OF INVENTION: DNA Molecules Encoding Plant
TITLE OF INVENTION: Protoporphyrinogen Oxidase and Inhibitor-Resistant Mutants
NUMBER OF SEQUENCES: 35
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 5939602artis Corporation
STREET: 520 White Plains Road, P.O. Box 2005
CITY: Tarrytown
STATE: NY
COUNTRY: USA
ZIP: 10591-9005
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/808,931
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/012,705
FILING DATE: 28-FEB-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/013,612
FILING DATE: 28-FEB-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/020,003
FILING DATE: 21-JUN-1996
ATTORNEY/AGENT INFORMATION:
NAME: Meigs, J. Timothy
REGISTRATION NUMBER: 38,241
REFERENCE/DOCKET NUMBER: CGC 1847
TELECOMMUNICATION INFORMATION:
TELEPHONE: (919) 541-8587
TELEFAX: (919) 541-8689
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 560 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-08-808-931-18

Query Match 85.7%; Score 18; DB 2; Length 560;
Best Local Similarity 66.7%; Pred. No. 2.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
Db 67 EAGSGS 72

RESULT 38
US-08-808-323-18
Sequence 18, Application US/08808323
Patent No. 6018105
GENERAL INFORMATION:
APPLICANT: Johnson, Marie
APPLICANT: Volrath, Sandra
APPLICANT: Ward, Eric
TITLE OF INVENTION: Promoters from Plant
TITLE OF INVENTION: Protoporphyrinogen Oxidase Genes
NUMBER OF SEQUENCES: 26
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 6018105artis Corporation
STREET: 520 White Plains Road, P.O. Box 2005
CITY: Tarrytown
STATE: NY
COUNTRY: USA
ZIP: 10591-9005

```

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/808,323
; FILING DATE:
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/012,705
; FILING DATE: 28-FEB-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/013,612
; FILING DATE: 28-FEB-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/020,003
; FILING DATE: 21-JUN-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Meigs, J. Timothy
; REGISTRATION NUMBER: 38,241
; REFERENCE/DOCKET NUMBER: CGC 1846
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (919) 541-8587
; TELEFAX: (919) 541-8689
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 560 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
US-08-808-323-18

```

```

Query Match      85.7%; Score 18; DB 3; Length 560;
Best Local Similarity 66.7%; Pred. NO. 2.le+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

```

Qy 1 eaqxss 6
Db 67 EAGSGS 72

```

```

RESULT 39
US-09-050-603A-18
; Sequence 18, Application US/09050603A
; Patent No. 6023012
; GENERAL INFORMATION:
; APPLICANT: Volrath, Sandra
; APPLICANT: Johnson, Marie
; APPLICANT: Potter, Sharon
; APPLICANT: Ward, Eric
; APPLICANT: Heifetz, Peter
; TITLE OF INVENTION: DNA Molecules Encoding Plant
; TITLE OF INVENTION: Protoporphyrinogen Oxidase
; NUMBER OF SEQUENCES: 37
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: No. 6023012artis Corporation
; STREET: 3054 Cornwallis Road
; CITY: Research Triangle Park
; STATE: NC
; COUNTRY: USA
; ZIP: 27709
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/050,603A
; FILING DATE: 30-MAR-1998
; CLASSIFICATION: 800

```

```

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/808,931
; FILING DATE: 28-FEB-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/012,705
; FILING DATE: 28-FEB-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/013,612
; FILING DATE: 28-FEB-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/020,003
; FILING DATE: 21-JUN-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Meigs, J. Timothy
; REGISTRATION NUMBER: 38,241
; REFERENCE/DOCKET NUMBER: CGC 1847
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (919) 541-8587
; TELEFAX: (919) 541-8689
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 560 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
US-09-050-603A-18

```

```

Query Match      85.7%; Score 18; DB 3; Length 560;
Best Local Similarity 66.7%; Pred. NO. 2.le+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

```

Qy 1 eaqxss 6
Db 67 EAGSGS 72

```

```

RESULT 40
US-09-102-420B-18
; Sequence 18, Application US/09102420B
; Patent No. 6084155
; GENERAL INFORMATION:
; APPLICANT: Volrath, Sandra
; APPLICANT: Johnson, Marie
; APPLICANT: Ward, Eric
; APPLICANT: Heifetz, Peter
; TITLE OF INVENTION: HERBICIDE-TOLERANT PROTOPORPHYRINOGEN
; TITLE OF INVENTION: OXIDASE ("PROTOX")
; NUMBER OF SEQUENCES: 43
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: No. 6084155artis Corporation
; STREET: 3054 Cornwallis Road
; CITY: Research Triangle Park
; STATE: NC
; COUNTRY: USA
; ZIP: 27709
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/102,420B
; FILING DATE: 22-JUN-1998
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 09/059,164
; FILING DATE: 13-APR-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 09/050,603
; FILING DATE: 30-MAR-1998
; PRIOR APPLICATION DATA:

```


; APPLICATION NUMBER: US 60/126,430
; FILING DATE: 11-MAR-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/808,931
; FILING DATE: 28-FEB-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/012,705
; FILING DATE: 28-FEB-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/013,612
; FILING DATE: 28-FEB-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/020,003
; FILING DATE: 21-JUN-1996
; APPLICATION NUMBER: US 08/472,028
; FILING DATE: 06-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Meigs, J. Timothy
; REGISTRATION NUMBER: 38,241
; REFERENCE/DOCKET NUMBER: CGC 1847/CIP3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (919) 541-8587
; TELEFAX: (919) 541-8689
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 560 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
; US-09-102-420B-18

Query Match 85.7%; Score 18; DB 3; Length 560;
Best Local Similarity 66.7%; Pred. No. 2.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 67 EAGSGS 72

RESULT 41
US-09-497-698-18
; Sequence 18, Application US/09497698
; Patent No. 6308458
; GENERAL INFORMATION:
; APPLICANT: Volrath, Sandra
; Johnson, Marie
; Ward, Eric
; Heifetz, Peter
; TITLE OF INVENTION: HERBICIDE-TOLERANT PROTOPORPHYRINOGEN
; OXIDASE ("PROTOX")
; NUMBER OF SEQUENCES: 43
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: No. 6308458artis Corporation
; STREET: 3054 Cornwallis Road
; CITY: Research Triangle Park
; STATE: NC
; COUNTRY: USA
; ZIP: 27709
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/497,698
; FILING DATE: 03-Feb-2000
; CLASSIFICATION: <Unknown>
; 30-MAR-1998
; 11-MAR-1998

; 28-FEB-1997
; 28-FEB-1996
; 28-FEB-1996
; 21-JUN-1996
; 06-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/102,420
; FILING DATE: <Unknown>
; APPLICATION NUMBER: US 09/050,603
; FILING DATE: 30-MAR-1998
; APPLICATION NUMBER: US 60/126,430
; FILING DATE: 11-MAR-1998
; APPLICATION NUMBER: US 08/808,931
; FILING DATE: 28-FEB-1997
; APPLICATION NUMBER: US 60/012,705
; FILING DATE: 28-FEB-1996
; APPLICATION NUMBER: US 60/013,612
; FILING DATE: 28-FEB-1996
; APPLICATION NUMBER: US 60/020,003
; FILING DATE: 21-JUN-1996
; APPLICATION NUMBER: US 08/472,028
; FILING DATE: 06-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Meigs, J. Timothy
; REGISTRATION NUMBER: 38,241
; REFERENCE/DOCKET NUMBER: CGC 1847/CIP3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (919) 541-8587
; TELEFAX: (919) 541-8689
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 560 amino acids
; TYPE: amino acid
; STRANDEDNESS: No. 6308458 Relevant
; TOPOLOGY: No. 6308458 Relevant
; MOLECULE TYPE: protein
; SEQUENCE DESCRIPTION: SEQ ID NO: 18:
US-09-497-698-18

Query Match 85.7%; Score 18; DB 4; Length 560;
Best Local Similarity 66.7%; Pred. No. 2.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 67 EAGSGS 72

RESULT 42
US-08-419-078-2
; Sequence 2, Application US/08419078
; Patent No. 5587306
; GENERAL INFORMATION:
; APPLICANT: HAWKINS, PHILLIP R.
; APPLICANT: SEILHAMER, JEFFREY J.
; TITLE OF INVENTION: PHOSPHOLIPASE C HOMOLOG
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3330 HILLVIEW AVENUE
; CITY: PALO ALTO
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/419,078
; FILING DATE:

CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: LUTHER, BARBARA J.
REGISTRATION NUMBER: 33954
REFERENCE/DOCKET NUMBER: PF0030 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-855-0572
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 566 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
IMMEDIATE SOURCE:
LIBRARY: No. 5587306e
CLONE: 9118
US-08-419-078-2

Query Match 85.7%; Score 18; DB 1; Length 566;
Best Local Similarity 66.7%; Pred. No. 2.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |
Db 338 EAGQS 343

RESULT 43
US-08-726-883-2
; Sequence 2, Application US/08726883
; Patent No. 5676946
; GENERAL INFORMATION:
; APPLICANT: HAWKINS, PHILLIP R.
; APPLICANT: SEILHAMER, JEFFREY J.
; TITLE OF INVENTION: PHOSPHOLIPASE C HOMOLOG
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3330 HILLVIEW AVENUE
; CITY: PALO ALTO
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/726.883
; FILING DATE: 04-OCT-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/419,078
; FILING DATE: 10-APR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: LUTHER, BARBARA J.
; REGISTRATION NUMBER: 33954
; REFERENCE/DOCKET NUMBER: PF0030 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-855-0572
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 566 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; IMMEDIATE SOURCE:

LIBRARY: No. 5676946e
CLONE: 9118
US-08-726-883-2

Query Match 85.7%; Score 18; DB 1; Length 566;
Best Local Similarity 66.7%; Pred. No. 2.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |
Db 338 EAGQS 343

RESULT 44
US-08-696-944-2
; Sequence 2, Application US/08696944
; Patent No. 5981831
; GENERAL INFORMATION:
; APPLICANT: Sumant CHENGAPPA
; APPLICANT: Susan A. HELLYER
; APPLICANT: John S. REID
; APPLICANT: Jacqueline DE SILVA
; TITLE OF INVENTION: No. 5981831el Exo-(1-4)-Beta-D Galactanase
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pillsbury Madison & Sutro, L.L.P.
; STREET: 1100 New York Avenue, N.W.
; CITY: Washington
; STATE: D.C.
; COUNTRY: U.S.A.
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5 inch disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: MS Word
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/696,944
; FILING DATE: 23-AUG-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB95/00372
; FILING DATE: 23-FEB-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: GB 9403423.8
; FILING DATE: 23-FEB-1994
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 730 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-696-944-2

Query Match 85.7%; Score 18; DB 2; Length 730;
Best Local Similarity 66.7%; Pred. No. 2.7e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |
Db 603 EAGSNS 608

RESULT 45
US-08-731-716-2
; Sequence 2, Application US/08731716
; Patent No. 5789202
; GENERAL INFORMATION:
; APPLICANT: Hoskins, JoAnn
; APPLICANT: Jaskunas, S. Richard
; APPLICANT: Rockey, Pamela K.
; APPLICANT: Zhao, Genshi

APPLICANT: Rosteck, Paul R. Jr.
APPLICANT: NO. 5789202ris, Franklin H.
TITLE OF INVENTION: Penicillin Binding Protein From
TITLE OF INVENTION: Streptococcus Pneumoniae
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Eli Lilly and Company
STREET: Lilly Corporate Center
CITY: Indianapolis
STATE: Indiana
COUNTRY: U.S.
ZIP: 46285
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/731,716
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Webster, Thomas D.
REGISTRATION NUMBER: 39,872
REFERENCE/DOCKET NUMBER: X-10,887
TELECOMMUNICATION INFORMATION:
TELEPHONE: 317-276-3334
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 731 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-731-716-2

Query Match 85.7%; Score 18; DB 1; Length 731;
Best Local Similarity 66.7%; Pred. No. 2.7e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 102 EAGALS 107

RESULT 46
US-09-651-656-19
Sequence 19, Application US/09651656
Patent No. 6340566
GENERAL INFORMATION:
APPLICANT: MCCUTHEN-MALONEY, SANDRA
APPLICANT: LAWRENCE LIVERMORE NATIONAL LABORATORY
TITLE OF INVENTION: DETECTION AND QUANTITATION OF SINGLE NUCLEOTIDE
POLYMORPHISMS, DNA SEQUENCE VARIATIONS, DNA MUTATIONS,
DNA DAMAGE AND DNA MISMATCHES
FILE REFERENCE: IL-10689
CURRENT APPLICATION NUMBER: US/09/651,656
CURRENT FILING DATE: 2000-08-29
PRIOR APPLICATION NUMBER: 60/192,764
PRIOR FILING DATE: 2000-03-28
NUMBER OF SEQ ID NOS: 106
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 19
LENGTH: 823
TYPE: PRT
ORGANISM: Homo sapiens
US-09-651-656-19

Query Match 85.7%; Score 18; DB 4; Length 823;
Best Local Similarity 66.7%; Pred. No. 3e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 321 EAGSGS 326

RESULT 47
US-08-434-730-14
Sequence 14, Application US/08434730
Patent No. 5637463
GENERAL INFORMATION:
APPLICANT: Dalton, Stephen
APPLICANT: Kochan, Jarema P
APPLICANT: Osborne, Mark A
TITLE OF INVENTION: METHOD TO DETECT PROTEIN-PROTEIN
INTERACTIONS
NUMBER OF SEQUENCES: 16
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hoffmann-La Roche Inc
STREET: 340 Kingsland Street
CITY: Nutley
STATE: NJ
COUNTRY: USA
ZIP: 07110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/434,730
FILING DATE: 04-MAY-1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Semionow, Raina
REGISTRATION NUMBER: 39022
REFERENCE/DOCKET NUMBER: 9069
TELECOMMUNICATION INFORMATION:
TELEPHONE: (201)235-4391
TELEFAX: (201)235-2363
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 968 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-08-434-730-14

Query Match 85.7%; Score 18; DB 1; Length 968;
Best Local Similarity 66.7%; Pred. No. 3.5e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 711 EAGVTS 716

RESULT 48
US-08-560-005-2
Sequence 2, Application US/08560005
Patent No. 6001354
GENERAL INFORMATION:
APPLICANT: Pot, David A.
APPLICANT: Williams, Lewis T.
APPLICANT: Jefferson, Anne Bennett
APPLICANT: Majerus, Philip W.
TITLE OF INVENTION: No. 6001354el Grb2 Associating Protein and Nucleic
Acids Encoding Therefor
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Crew
STREET: One Market Plaza, Steuart Tower, Suite 2000

```

; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/560,005
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Dow, Karen B.
; REGISTRATION NUMBER: 29,684
; REFERENCE/DOCKET NUMBER: 2307K-0624000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-326-2400
; TELEFAX: 415-326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 976 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-560-005-2

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Query Match 85.7%; Score 18; DB 3; Length 976;
 Best Local Similarity 66.7%; Pred. No. 3.6e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

Qy 1 eagxxs 6
    ||| |
Db 496 EAGVTS 501

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```

RESULT 49
US-09-195-868-14
; Sequence 14, Application US/09195868
; Patent No. 6090621
; GENERAL INFORMATION:
; APPLICANT: KAVANAUGH MD, MICHAEL
; APPLICANT: POT PH.D., DAVID
; APPLICANT: WILLIAMS MDPHD, LEWIS T.
; TITLE OF INVENTION: SIGNALING INOSITOL POLYPHOSPHATE
; TITLE OF INVENTION: 5-PHOSPHATASES (SIPS)
; NUMBER OF SEQUENCES: 33
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: CHIRON CORPORATION
; STREET: 4560 HORTON STREET
; CITY: EMERYVILLE
; STATE: CA
; COUNTRY: USA
; ZIP: 94608
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/195,868
; FILING DATE:
; CLASSIFICATION:
; APPLICATION NUMBER: US/09/195,868
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: FIRESTONE, LEIGH H.
; REGISTRATION NUMBER: 36,831
; REFERENCE/DOCKET NUMBER: 1182.004
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 510-923-2707

```

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; TELEFAX: 510-655-3542
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 976 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-195-868-14

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Query Match 85.7%; Score 18; DB 3; Length 976;
 Best Local Similarity 66.7%; Pred. No. 3.6e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Qy 1 eagxxs 6
    ||| |
Db 496 EAGVTS 501

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RESULT 50
US-09-418-540-2
; Sequence 2, Application US/09418540
; Patent No. 6296848
; GENERAL INFORMATION:
; APPLICANT: Pot, David A.
; APPLICANT: Williams, Lewis T.
; APPLICANT: Jefferson, Anne Bennett
; APPLICANT: Majerus, Philip W.
; TITLE OF INVENTION: No. 6296848el Grb2 Associating Protein and Nucleic
; TITLE OF INVENTION: Acids Encoding Therefor
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew
; STREET: One Market Plaza, Steuart Tower, Suite 2000
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94105
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/418,540
; FILING DATE: 14-OCT-1999
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/560,005
; FILING DATE: 17-NOV-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Dow, Karen B.
; REGISTRATION NUMBER: 29,684
; REFERENCE/DOCKET NUMBER: 2307K-0624000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-326-2400
; TELEFAX: 415-326-2422
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 976 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-418-540-2

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Query Match 85.7%; Score 18; DB 4; Length 976;
 Best Local Similarity 66.7%; Pred. No. 3.6e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Qy 1 eagxxs 6
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Db 496 EAGVTS 501

Search completed: August 30, 2002, 15:04:45
Job time: 8556 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: August 30, 2002, 15:00:04 ; Search time 26.93 seconds
(without alignments)
21.409 Million cell updates/sec

Title: BASK-853-CLAIM4

Perfect score: 21
Sequence: 1 eagxxs 6

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283138 seqs, 96089334 residues

Total number of hits satisfying chosen parameters: 283138

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 50 summaries

Database : PIR_71.*

1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19	90.5	37	2 A48620	adhesin - Staphylo
2	19	90.5	83	2 AF2563	hypothetical prote
3	19	90.5	122	2 C84320	hypothetical prote
4	19	90.5	143	2 B72627	hypothetical prote
5	19	90.5	145	2 F82189	hypothetical prote
6	19	90.5	164	2 T11215	hypothetical prote
7	19	90.5	200	2 T29807	hypothetical prote
8	19	90.5	201	2 T23855	hypothetical prote
9	19	90.5	204	2 A70844	probable moe3 pro
10	19	90.5	212	2 AD1560	two-component resp
11	19	90.5	212	2 AF1202	two-component resp
12	19	90.5	220	2 AH0459	Sec-independent pr
13	19	90.5	245	2 E84169	hypothetical prote
14	19	90.5	246	2 AE1029	probable exported
15	19	90.5	257	2 T48058	RING-H2 zinc finge
16	19	90.5	260	2 B90026	hypothetical prote
17	19	90.5	263	2 G87721	protein ZC123.3 (i
18	19	90.5	266	2 JC1071	coat protein - soy
19	19	90.5	267	2 S18931	coat protein - soy
20	19	90.5	302	2 H82638	hypothetical prote
21	19	90.5	303	1 D64070	ATP phosphoribosyl
22	19	90.5	311	2 A56235	transcription acti
23	19	90.5	322	2 S38091	hypothetical prote
24	19	90.5	323	2 I49529	transcription fact
25	19	90.5	332	2 A38873	myristylated alani
26	19	90.5	352	2 S16547	neutral adenylase
27	19	90.5	352	2 G95872	probable adenylate
28	19	90.5	357	2 T01571	hypothetical prote
29	19	90.5	363	2 A83177	probable N-acetylgl

ALIGNMENTS

RESULT 1

A48620 adhesin - Staphylococcus aureus (fragment)

C:Species: Staphylococcus aureus

C:Date: 07-Apr-1994 #sequence_revision 18-Nov-1994 #text_change 03-Mar-1995

C:Accession: A48620

R:Patti, J.M.; Boles, J.O.; Hook, M.

Biochemistry 32:11428-11435, 1993

A:Title: Identification and biochemical characterization of the ligand binding domain

A:Reference number: A48620; MUID:94032261

A:Contents: FDA 574

A:Accession: A48620

A>Status: preliminary; not compared with conceptual translation

A:Molecule type: nucleic acid; protein

A:Residues: 1-37 <PAI>

A>Note: sequence extracted from NCBI backbone (NCBIP:138726)

Query Match 90.5%; Score 19; DB 2; Length 37;
Best Local Similarity 66.7%; Pred.No. 89;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

||| |

Db 16 EAGTSS 21

RESULT 2

AF2563 hypothetical protein asl8505 [imported] - Anabaena sp. (strain PCC 7120) plasmid pcc7

C:Species: Anabaena sp.

A>Note: Anabaena sp. (strain PCC 7120) is a synonym of Nostoc sp. strain PCC 7120

C:Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 11-Jan-2002

C:Accession: AF2563

R:Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Irigu

Nakazaki, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Tabata

DNA Res. 8, 205-213, 2001

A:Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium

A:Reference number: AB1807; MUID:21595285; PMID:11759840

A:Accession: AF2563

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-83 <KUR>

A:Cross-references: GB:AP003604; PIDN:BAE77424.1; PID:g17134868; GSPDB:GN00183

A:Experimental source: strain PCC 7120

C:Genetics:

A:Gene: asl8505

A:Genome: plasmid

Query Match 90.5%; Score 19; DB 2; Length 83;
Best Local Similarity 66.7%; Pred. No. 1.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |

Db 78 EAGASS 83

RESULT 3

C84320 hypothetical protein Vngl678h [imported] - Halobacterium sp. NRC-1

C:Species: Halobacterium sp. NRC-1

C:Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001

C:Accession: C84320
R;Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky, S.;
Leithauser, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Jablon
Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000

A:Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.; Li
A:Title: Genome sequence of Halobacterium species NRC-1.

A:Reference number: A84160; MUID:20504483

A:Accession: C84320

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-122 <STO>

A:Cross-references: GB:AE004437; NID:g10581147; PIDN:AAG19927.1; GSPDB:GN00138

C:Genetics:

A:Gene: VNGl678H

Query Match 90.5%; Score 19; DB 2; Length 122;
Best Local Similarity 66.7%; Pred. No. 2.8e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |

Db 57 EAGASS 62

RESULT 4

B72627 hypothetical protein APEI474 - Aeropyrum pernix (strain K1)

C:Species: Aeropyrum pernix

C:Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Aug-1999

C:Accession: B72627
R;Kawarayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Tanaka, Y.; Jin-no, K.; Takah
awa, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, Y.; Kudoh, Y.; Yamazaki, J.; K
DNA Res. 6, 83-101, 1999

A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyr

A:Reference number: A72450; MUID:99310339

A:Accession: B72627

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-143 <KAW>

A:Cross-references: DDBJ:AP000061; NID:g5104821; PIDN:BAA80472.1; PID:g1044258; PID:g510

C:Genetics:

A:Gene: APEI474

Query Match 90.5%; Score 19; DB 2; Length 143;
Best Local Similarity 66.7%; Pred. No. 3.2e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |

Db 85 EAGAAS 90

RESULT 5

F82189

hypothetical protein VC1536 [imported] - Vibrio cholerae (strain N16961 serogroup O1)

C:Species: Vibrio cholerae

C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Feb-2001

C:Accession: F82189
R;Heidelberg, J.F.; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.; Dodson, R.
chardson, D.; Ermolaeva, M.D.; Vamathevan, J.; Bass, S.; Qlin, H.; Dragoi, I.; Sellers
1, R.R.; Mekalanos, J.J.; Venter, J.C.; Fraser, C.M.
Nature 406, 477-483, 2000

A:Title: DNA Sequence of both chromosomes of the cholera pathogen Vibrio cholerae.

A:Reference number: A82035; MUID:20406833

A:Accession: F82189

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-145 <HEI>

A:Cross-references: GB:AF004231; GB:AE003852; NID:g9656027; PIDN:AAF94690.1; GSPDB:GN

A:Experimental source: serogroup O1; strain N16961; biotype El Tor

C:Genetics:

A:Gene: VC1536

A:Map position: 1

Query Match 90.5%; Score 19; DB 2; Length 145;
Best Local Similarity 66.7%; Pred. No. 3.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |

Db 114 EAGSTS 119

RESULT 6

T11215

hypothetical protein 5 - Streptomyces glaucescens

C:Species: Streptomyces glaucescens

C:Date: 16-Jul-1999 #sequence_revision 16-Jul-1999 #text_change 16-Jul-1999

C:Accession: T11215

R;Summers, R.G.; Ali, A.; Shen, B.; Wessel, W.A.; Hutchinson, C.R.

Biochemistry 34, 9389-9402, 1995

A:Title: Malonyl-coenzyme A:acyl carrier protein acyltransferase of Streptomyces glau

A:Reference number: 217254; MUID:95352622

A:Accession: T11215

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-164 <SUM>

A:Cross-references: EMBL:L43074; NID:g870805; PID:g870810

Query Match 90.5%; Score 19; DB 2; Length 164;
Best Local Similarity 66.7%; Pred. No. 3.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
||| |

Db 55 EAGTAS 60

RESULT 7

T29807

hypothetical protein C25A8.2 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999

C:Accession: T29807

R;Latreille, P.; Stellyes, L.

submitted to the EMBL Data Library, June 1996

A:Description: The sequence of C. elegans cosmid C25A8.

A:Reference number: 220689

A:Accession: T29807

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-200 <LAT>

A:Cross-references: EMBL:U61958; PIDN:AA803180.1; GSPDB:GN00022; CESP:C25A8.2

A:Experimental source: strain Bristol N2; clone C25A8
C:Genetics:
A:Gene: CESP:C25A8.2
A:Map position: 4
A:Introns: 173/3

Query Match 90.5%; Score 19; DB 2; Length 200;
Best Local Similarity 66.7%; Pred. No. 4.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 109 EGAAS 114

RESULT 8
T23855
hypothetical protein R02D5.7 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
C:Accession: T23855
R:Matthews, L.
submitted to the EMBL Data Library, August 1996
A:Reference number: Z19808
A:Accession: T23855
A:Status: preliminary; translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 1-201 <WIL>
A:Cross-references: EMBL:278015; PIDN:CAB01436.1; GSPDB:GN00023; CESP:R02D5.7
A:Experimental source: clone R02D5
C:Genetics:
A:Gene: CESP:R02D5.7
A:Map position: 5
A:Introns: 174/3

Query Match 90.5%; Score 19; DB 2; Length 201;
Best Local Similarity 66.7%; Pred. No. 4.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 110 EGAAS 115

RESULT 9
A70844
probable moaE3 protein - Mycobacterium tuberculosis (strain H37RV)
C:Species: Mycobacterium tuberculosis
C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999
C:Accession: A70844
R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.; Connor, R.; Davies, R.; Devlin, K.; Felwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.; Nature 393, 537-544, 1998
A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome
A:Reference number: A70500; MUID:98295987
A:Accession: A70844
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-204 <COL>
A:Cross-references: GB:AL021841; GB:AL123456; NID:g3261517; PIDN:CAA17094.1; PID:el25115
A:Experimental source: strain H37RV
C:Genetics:
A:Gene: moaE3

Query Match 90.5%; Score 19; DB 2; Length 204;
Best Local Similarity 66.7%; Pred. No. 4.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 153 EAGTAS 158

RESULT 10

AD1560
two-component response regulator, in particular B. subtilis YvqC protein homolog lin1
C:Species: Listeria innocua
C:Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 14-Dec-2001
C:Accession: AD1560
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloec
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi,
D.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.; Maitournam, A.;
ok, C.; Schlueter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehla
A:Title: Comparative genomics of Listeria species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669

A:Accession: AD1560
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-212 <GLA>
A:Cross-references: GB:AL592022; PIDN:CAC96252.1; PID:gl6413480; GSPDB:GN00178
A:Experimental source: strain Clp11262
C:Genetics:
A:Gene: lin1021
C:Superfamily: regulatory protein comA; response regulator homology

Query Match 90.5%; Score 19; DB 2; Length 212;
Best Local Similarity 66.7%; Pred. No. 4.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 95 EAGASS 100

RESULT 11

AF1202
two-component response regulator, in particular B. subtilis YvqC protein homolog lmo1
C:Species: Listeria monocytogenes
C:Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 14-Dec-2001
C:Accession: AF1202
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloec
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi,
D.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.; Maitournam, A.;
ok, C.; Schlueter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehla
A:Title: Comparative genomics of Listeria species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669

A:Accession: AF1202
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-212 <GLA>
A:Cross-references: GB:NC_003210; PIDN:CAC99100.1; PID:gl6410424; GSPDB:GN00177
A:Experimental source: strain EGD-e
C:Genetics:
A:Gene: lmo1022
C:Superfamily: regulatory protein comA; response regulator homology

Query Match 90.5%; Score 19; DB 2; Length 212;
Best Local Similarity 66.7%; Pred. No. 4.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 95 EAGASS 100

RESULT 12

AH0459
Sec-Independent protein translocase protein TatB [imported] - Yersinia pestis (strain CO
C:Species: Yersinia pestis
C:Date: 02-Nov-2001 #sequence_revision 02-Nov-2001 #text_change 02-Nov-2001
C:Accession: AH0459
R:Parkhill, J.; Wren, B.W.; Thomson, N.R.; Titball, R.W.; Holden, M.T.G.; Prentice, M.B.
deno-Tarraga, A.M.; Chillingworth, T.; Cronin, A.; Davies, R.M.; Davis, P.; Dougan, G.;
Il, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; Whitehead, S.; Barrell,
Nature 413, 523-527, 2001
A:Title: Genome sequence of Yersinia pestis, the causative agent of plague.
A:Reference number: AB0001; MUID:21470413; PMID:11586360
A:Accession: AH0459
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-220 <KUR>
A:CROSS-references: GB:AL590842; PIDN:CAC93244.1; PID:g15981690; GSPDB:GN00175
C:Genetics:
A:Gene: tatB

Query Match 90.5%; Score 19; DB 2; Length 220;
Best Local Similarity 66.7%; Pred. No. 4.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 143 EAGTAS 148

RESULT 13

B84169
hypothetical protein pimT1 [imported] - Halobacterium sp. NRC-1
C:Species: Halobacterium sp. NRC-1
C:Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
C:Accession: B84169
R:Ng, W.V.; Kennedy, S.P.; Mahairas, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky, S.
Leithauser, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Jabid
Jung, K.H.; Alam, M.; Freitas, T.
Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000
A:Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ebhardt, H.; Lowe, T.M.; Li
A:Title: Genome sequence of Halobacterium species NRC-1.
A:Reference number: B84160; MUID:20504483
A:Accession: B84169
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-245 <STO>
A:CROSS-references: GB:AE004437; MID:g10579741; PIDN:AAG18721.1; GSPDB:GN00138
C:Genetics:
A:Gene: pimT1

Query Match 90.5%; Score 19; DB 2; Length 245;
Best Local Similarity 66.7%; Pred. No. 5.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 151 EAGAAS 156

RESULT 14

AE1029
probable exported protein STY4558 [imported] - Salmonella enterica subsp. enterica serov
C:Species: Salmonella enterica subsp. enterica serovar Typhi
A:Note: this species has also been called Salmonella typhi
C:Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 09-Nov-2001
C:Accession: AE1029
R:Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; Wain, J.; Churcher,
th, T.; Connor, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.; White, N.; Farrar,
S.; Moulé, S.; O'Gaora, P.
Nature 413, 848-852, 2001
A:Authors: Parry, C.; Quall, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.

A:Title: Complete genome sequence of a multiple drug resistant Salmonella enterica se
A:Reference number: AB0502; PMID:11677608
A:Accession: AE1029
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-246 <PAR>
A:CROSS-references: GB:AL513382; PIDN:CAD09334.1; PID:g16505334; GSPDB:GN00176
C:Genetics:
A:Gene: STY4558

Query Match 90.5%; Score 19; DB 2; Length 246;
Best Local Similarity 66.7%; Pred. No. 5.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 93 EAGSAS 98

RESULT 15

T48058
RING-H2 zinc finger protein ATL5 - Arabidopsis thaliana
N:Alternate names: protein F26K9.120
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 20-Apr-2000 #sequence_revision 20-Apr-2000 #text_change 08-Dec-2000
C:Accession: T48058
R:Bloeker, H.; Mewes, H.W.; Rudd, S.; Lemcke, K.; Mayer, K.F.X.; Quetier, F.; Salano
submitted to the Protein Sequence Database, March 2000
A:Reference number: Z24465
A:Accession: T48058
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-257 <BLO>
A:CROSS-references: EMBL:AL162651
A:Experimental source: cultivar Columbia; BAC clone F26K9
C:Genetics:
A:Map position: 3
A:Note: F26K9.120
C:Superfamily: Arabidopsis hypothetical protein F19T3.22; RING finger homology
F:109-160/Domain: RING finger homology <RRN>

Query Match 90.5%; Score 19; DB 2; Length 257;
Best Local Similarity 66.7%; Pred. No. 5.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 184 EAGSSS 189

RESULT 16

B90026
hypothetical protein modA [imported] - Staphylococcus aureus (strain N315)
C:Species: Staphylococcus aureus
C:Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 22-Oct-2001
C:Accession: B90026
R:Kuroda, M.; Ohta, T.; Uchiyama, I.; Baba, T.; Yuzawa, H.; Kobayashi, I.; Cui, L.; O
ma, A.; Mizutani-Ui, Y.; Kobayashi, N.; Sawano, T.; Inoue, R.; Kaito, C.; Sekimizu, K.
C.; Shiba, T.; Hattori, M.; Ogasawara, N.; Hayashi, H.; Hiramatsu, K.
Lancet 357, 1225-1240, 2001
A:Title: Whole genome sequencing of methicillin-resistant Staphylococcus aureus.

A:Reference number: A89758; MUID:21311952; PMID:11418146
A:Accession: B90026
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-260 <KUR>

A:CROSS-references: GB:BA000018; PID:g13702079; PIDN:BA043371.1; GSPDB:GN00149
A:Experimental source: strain N315
C:Genetics:
A:Gene: modA
C:Superfamily: molybdate-binding periplasmic protein

Query Match 90.5%; Score 19; DB 2; Length 260;
 Best Local Similarity 66.7%; Pred. No. 5.7e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 227 EAGATS 232

RESULT 17

G87721
 protein ZC123.3 [imported] - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 10-May-2001
 C:Accession: G87721
 R:Anonymous, The C. elegans Sequencing Consortium.
 Science 282, 2012-2018, 1998
 A:Title: Genome sequence of the nematode C. elegans: a platform for investigating biological processes
 A:Reference number: A75000; MUID:99069613; PMID:9851916
 A:Note: see websites genome.wustl.edu/gsc/C.elegans/ and www.sanger.ac.uk/Projects/C_elegans/
 A:Note: published errata appeared in Science 283, 35, 1999; Science 283, 2103, 1999; and Science 283, 2103, 1999
 A:Accession: G87721
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-263 <STO>
 A:Cross-references: GB:chr.I; PIDN:AAB97603.1; PID:g2804499; GSPDB:GN00019; CESP:ZC123.3
 A:Note: contains similarity to C2H2-type zinc fingers
 C:Genetics:
 A:Gene: ZC123.3
 A:Map position: 1

Query Match 90.5%; Score 19; DB 2; Length 263;
 Best Local Similarity 66.7%; Pred. No. 5.8e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 31 EAGASS 36

RESULT 18

JC1071
 coat protein - soybean mosaic virus
 C:Species: soybean mosaic virus, SBMV
 C:Date: 02-Aug-1995 #sequence_revision 19-Oct-1995 #text_change 17-Nov-2000
 C:Accession: JC1071
 R:Chu, R.Y.; Leng, X.H.; Bao, Y.M.; Pan, N.S.; Pu, Z.Q.
 Acta Bot. Sin. 34, 523-528, 1992
 A:Title: Amplification of soybean mosaic virus coat protein gene by polymerase chain reaction
 A:Reference number: JC1071
 A:Accession: JC1071
 A:Molecule type: DNA
 A:Residues: 1-266 <CHU>
 C:Superfamily: tobacco etch virus genome polyprotein

Query Match 90.5%; Score 19; DB 2; Length 266;
 Best Local Similarity 66.7%; Pred. No. 5.8e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 25 EAGTSS 30

RESULT 19

S18931
 coat protein - soybean mosaic virus (fragment)
 C:Species: soybean mosaic virus, SBMV
 C:Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 23-Mar-2001

C:Accession: S18931
 R:Chu, R.

Submitted to the EMBL Data Library, January 1992
 A:Description: CDNA sequence of the gene encoding coat protein of SBMV.
 A:Reference number: S18931

A:Accession: S18931
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-267 <CHU>
 A:Cross-references: EMBL:X63771; NID:g61983; PIDN:CAA45307.1; PID:g61984
 C:Superfamily: tobacco etch virus genome polyprotein

Query Match 90.5%; Score 19; DB 2; Length 267;
 Best Local Similarity 66.7%; Pred. No. 5.8e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 26 EAGTSS 31

RESULT 20

H82638
 hypothetical protein XF1783 [imported] - Xylella fastidiosa (strain 9a5c)
 C:Species: Xylella fastidiosa
 C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 20-Aug-2000
 C:Accession: H82638
 R:Anonymous, The Xylella fastidiosa Consortium of the Organization for Nucleotide Sequencing
 Nature 406, 151-157, 2000
 A:Title: The genome sequence of the plant pathogen Xylella fastidiosa.
 A:Reference number: A82515; MUID:20365717
 A:Note: for a complete list of authors see reference number A59328 below
 A:Accession: H82638
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-302 <SIM>
 A:Experimental source: strain 9a5c
 A:Cross-references: GB:AE004000; GB:AE003849; NID:g9106850; PIDN:AAF84591.1; GSPDB:GN
 R:Simpson, A.J.G.; Reinach, F.C.; Arruda, P.; Abreu, F.A.; Acencio, M.; Alvarenga, R.
 Briones, M.R.S.; Bueno, M.R.P.; Camargo, A.A.; Camargo, D.M.; Carraro, D.M.; Carrier
 as-Neto, E.; Docena, C.; El-Dorri, H.; Facincani, A.P.; Ferreira, A.J.S.
 Submitted to GenBank, June 2000
 A:Authors: Ferreira, V.C.A.; Ferro, J.A.; Fraga, J.S.; Franca, S.C.; Franco, M.C.; Fr
 J.D.; Junqueira, M.L.; Kemper, E.L.; Kitajima, J.P.; Krieger, J.E.; Kuramae, E.E.; La
 chado, M.A.; Madeira, A.M.B.N.; Madeira, H.M.F.; Marino, C.L.; Marques, M.V.; Martins
 A:Authors: Martins, E.M.F.; Matsukuma, A.Y.; Menck, C.F.M.; Miracca, E.C.; Miyaki, C.
 F.G.; Nunes, L.R.; Oliveira, M.A.; de Oliveira, M.C.; de Oliveira, R.C.; Palmieri, R.
 Rodrigues, V.; Rosa, A.J. de M.; de Rosa Jr., V.E.; de Sa, R.G.; Santelli, R.V.; Sawa
 A:Authors: da Silva, A.C.R.; da Silva, F.R.; da Silva, A.M.; Silva Jr., W.A.; da Silv
 M.; Tshukako, M.H.; Vallada, H.; Van Sluys, M.A.; Verjovski-Almeida, S.; Vettore, A.L.
 A:Reference number: A59328
 A:Contents: annotation
 C:Genetics:
 A:Gene: XF1783

Query Match 90.5%; Score 19; DB 2; Length 302;
 Best Local Similarity 66.7%; Pred. No. 6.6e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
 ||| |
 Db 106 EAGTAS 111

RESULT 21

D64070
 ATP phosphoribosyltransferase (EC 2.4.2.17) - Haemophilus influenzae (strain Rd KW20)
 C:Species: Haemophilus influenzae
 C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 17-Mar-2000
 C:Accession: D64070
 R:Fleischmann, R.D.; Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage

; Gocayne, J.D.; Scott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.M.; Weidman, J.
D.M.; Brandon, R.C.; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Geoghagen, N.S.M.
Science 269, 496-512, 1995
A:Authors: Gnehm, C.L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter,
A.:Title: Whole-genome random sequencing and assembly of Haemophilus influenzae Rd.
A:Reference number: A64000; MUID:95350630
A:Accession: D64070
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-303 <TIGR>
A:Cross-references: GB:U32729; GB:L42023; NID:g1573439; PIDN:AAC22127.1; PID:g1573446; T
A:Note: named as homolog to a protein from Escherichia coli
C:Superfamily: ATP phosphoribosyltransferase; ATP phosphoribosyltransferase homology
C:Keywords: glycosyltransferase; histidine biosynthesis; pentosyltransferase

Query Match 90.5%; Score 19; DB 1; Length 303;
Best Local Similarity 66.7%; Pred. No. 6.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 eagxxs 6
||| |
Db 287 EAGASS 292

RESULT 22
A56235
transcription activator MafB - chicken
C:Species: Gallus gallus (chicken)
C:Date: 03-Oct-1995 #sequence_revision 03-Oct-1995 #text_change 20-Jun-2000
C:Accession: A56235
R:Kataoka, K.; Fujiwara, K.T.; Noda, M.; Nishizawa, M.
Mol. Cell. Biol. 14, 7581-7591, 1994
A:Title: MafB, a new Maf family transcription activator that can associate with Maf and
A:Reference number: A56235; MUID:95021288
A:Accession: A56235
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-311 <KAT>
A:Cross-references: GB:D28600; NID:g516723; PIDN:BAA05938.1; PID:g516724
C:Genetics:
A:Introns: #status absent
C:Superfamily: maf transforming protein; maf homology
C:Keywords: DNA binding; homodimer; leucine zipper
F:200-289/Domain: maf homology <MAF>

Query Match 90.5%; Score 19; DB 2; Length 311;
Best Local Similarity 66.7%; Pred. No. 6.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 eagxxs 6
||| |
Db 296 EAGSTS 301

RESULT 23
S38091
hypothetical protein YKR022c - yeast (Saccharomyces cerevisiae)
C:Species: Saccharomyces cerevisiae
C:Date: 03-May-1994 #sequence_revision 03-May-1994 #text_change 29-Oct-1999
C:Accession: S38091
R:Duesterhoeft, A.; Moestl, D.; Pohlmann, R.; Philippsen, P.
submitted to the Protein Sequence Database, March 1994
A:Reference number: S37811
A:Accession: S38091
A:Molecule type: DNA
A:Residues: 1-322 <DUE>
A:Cross-references: EMBL:Z28247; NID:g486444; PID:g486445; GSPDB:GN00011; MIPS:YKR022c
A:Experimental source: strain S288C
C:Genetics:
A:Gene: MIPS:YKR022c
A:Map position: 11R

Query Match 90.5%; Score 19; DB 2; Length 322;
Best Local Similarity 66.7%; Pred. No. 7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 eagxxs 6
||| |
Db 119 EAGSSS 124

RESULT 24
I49529
transcription factor-kr - mouse
C:Species: Mus musculus (house mouse)
C:Date: 09-Mar-1996 #sequence_revision 09-Mar-1996 #text_change 16-Jul-1999
C:Accession: I49529
R:Cordes, S.P.; Barsh, G.S.
Cell 79, 1025-1034, 1994
A:Title: The mouse segmentation gene kr encodes a novel basic domain-leucine zipper t
A:Reference number: A55200; MUID:95094266
A:Accession: I49529
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-323 <RES>
A:Cross-references: GB:L36435; NID:g625043; PIDN:AAA65689.1; PID:g625044
C:Superfamily: maf transforming protein; maf homology
C:Keywords: leucine zipper; transcription factor
F:212-301/Domain: maf homology <MAF>

Query Match 90.5%; Score 19; DB 2; Length 323;
Best Local Similarity 66.7%; Pred. No. 7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 eagxxs 6
||| |
Db 308 EAGSTS 313

RESULT 25
A38873
myristylated alanine-rich protein kinase C substrate - human
N:Alternate names: acidic calmodulin-binding 80K protein; MARCKS
C:Species: Homo sapiens (man)
C:Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change 20-Jun-2000
C:Accession: A38873; A42977; A40758; S29269
R:Shimizu, N.
submitted to DBJ, September 1991
A:Reference number: A38873
A:Accession: A38873
A:Molecule type: mRNA
A:Residues: 1-332 <SHI>
A:Cross-references: GB:D10522; GB:D90498; NID:g219893; PIDN:BAA01392.1; PID:g219894
R:Sakai, K.; Hirai, M.; Kudoh, J.; Minoshima, S.; Shimizu, N.
Genomics 14, 175-178, 1992
A:Title: Molecular cloning and chromosomal mapping of a cDNA encoding human 80K-L pro
A:Reference number: A42977; MUID:93052291
A:Accession: A42977
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-13, 'G', 15-332 <SAK>
A:Cross-references: GB:D90498
A:Experimental source: squamous carcinoma cells A431
A:Note: sequence extracted from NCBI backbone (NCBIP:118653)
R:Harlan, D.M.; Graff, J.M.; Stumpo, D.J.; Eddy Jr., R.L.; Shows, T.B.; Boyle, J.M.;
J. Biol. Chem. 266, 14399-14405, 1991
A:Title: The human myristoylated alanine-rich C kinase substrate (MARCKS) gene (MACS)
A:Reference number: A40758; MUID:91317795
A:Accession: A40758
A:Molecule type: mRNA
A:Residues: 1-83, 'A', 85-118, 'P', 120-233, 'W', 235-286, 'LVC', 290, 'RRGSGPRGGARRSLNQ', 30
A:Cross-references: GB:M68956

A:Note: the authors translated the codon GGC for residue 53 as Arg
R:Herget, T.; Brooks, S.F.; Broad, S.; Rozenfurt, E.
Eur. J. Biochem. 209, 7-14, 1992
A:Title: Relationship between the major protein kinase C substrates acidic 80-kDa protein
or equivalent genes in different species.
A:Reference number: S29267; PMID:93011168

A:Accession: S29269
A:Molecule type: mRNA
A:Residues: 189-223,'R',225-234,'E',236-322 <HER>
A:Comment: This protein is a major cellular substrate for protein kinase C and plays a role
C:Comment: It binds to calmodulin in one to one molar ratio in the presence of calcium and
C:Genetics:
A:Gene: GDB:MACS

A:Cross-references: GDB:118835; OMIM:177061
A:Map position: 6q22.2-6q22.2
C:Superfamily: neurofilament triplet H protein
C:Keywords: actin binding; blocked amino end; calmodulin binding; lipoprotein; myristyla
F:2/Modified site: myristylated amino end (Gly) (in mature form) #status predicted
F:159,163,167,170/Binding site: phosphate (Ser) (covalent) (by protein kinase C) #status

Query Match 90.5%; Score 19; DB 2; Length 332;
Best Local Similarity 66.7%; Pred. No. 7.2e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 206 EAGAAS 211

RESULT 26

S16547
neutral proteinase II - Aspergillus oryzae
C:Species: Aspergillus oryzae
C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 01-Dec-2000
C:Accession: S16547; S47562
R:Atsumi, H.; Murakami, S.; Tsuji, R.F.; Ishida, Y.; Murakami, K.; Masaki, A.; Kawabe,
Mol. Gen. Genet. 228, 97-103, 1991
A:Title: Cloning and expression in yeast of a cDNA clone encoding Aspergillus oryzae neu
A:Reference number: S16547; PMID:91360097
A:Accession: S16547
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-352 <TAT>
R:Atsumi, H.; Ikegaya, K.; Murakami, S.; Kawabe, H.; Nakano, E.; Motai, H.
Biochim. Biophys. Acta 1208, 179-185, 1994
A:Title: Elucidation of the thermal stability of the neutral proteinase II from Aspergil
A:Reference number: S47562; PMID:94368822
A:Accession: S47562
A:Status: preliminary
A:Molecule type: protein
A:Residues: 176-352 <TAA>
C:Superfamily: Penicillium citrinum penicillolysin

Query Match 90.5%; Score 19; DB 2; Length 352;
Best Local Similarity 66.7%; Pred. No. 7.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 240 EAGSTS 245

RESULT 27

G95872
probable adenylate cyclase protein [imported] - Sinorhizobium meliloti (strain 1021) mag
C:Species: Sinorhizobium meliloti
C:Date: 24-Aug-2001 #sequence_revision 24-Aug-2001 #text_change 30-Sep-2001
C:Accession: G95872
R:Finan, T.M.; Weidner, S.; Wong, K.; Buhrmester, J.; Chain, P.; Vorholter, F.J.; Hernan
Proc. Natl. Acad. Sci. U.S.A. 98, 9889-9894, 2001

A:Title: The complete sequence of the 1.683-kb pSymb megaplasmid from the N2-fixing e
A:Reference number: A95842; PMID:21396508; PMID:11481431
A:Accession: G95872
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-352 <KUR>
A:Cross-references: GB:AL591985; PIDN:CAC48647.1; PID:gi15140119; GSPDB:GN00167
A:Experimental source: strain 1021, megaplasmid pSymb
R:Galibert, F.; Finan, T.M.; Long, S.R.; Puhler, A.; Abola, P.; Ampe, F.; Barloy-Hubl
pella, D.; Chain, P.; Cowie, A.; Davis, R.W.; Dreano, S.; Federspiel, N.A.; Fisher, R.
L.; Hyman, R.W.; Jones, T.
Science 293, 668-672, 2001

A:Authors: Kahn, D.; Kahn, M.L.; Kalman, S.; Keating, D.H.; Kiss, E.; Komp, C.; Lelau
hebaull, P.; Vandenberg, M.; Vorholter, F.J.; Weidner, S.; Wells, D.H.; Wong, K.; Yeh,
A:Title: The composite genome of the legume symbiont Sinorhizobium meliloti.
A:Reference number: A96039; PMID:21368234; PMID:11474104
A:Contents: annotation
C:Genetics:
A:Gene: SMD20257
A:Genome: plasmid

Query Match 90.5%; Score 19; DB 2; Length 352;
Best Local Similarity 66.7%; Pred. No. 7.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 13 EAGTSS 18

RESULT 28

T01571
hypothetical protein A_TM018A10.10 - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 19-Feb-1999 #sequence_revision 19-Feb-1999 #text_change 24-Nov-1999
C:Accession: T01571
R:Dempsey, S.; Harper, M.
Submitted to the EMBL Data Library, July 1997
A:Description: The sequence of A. thaliana TM018A10.
A:Reference number: Z14348
A:Accession: T01571
A:Status: translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-357 <DEM>
A:Cross-references: EMBL:AF013294; NID:g2252848; PID:g2252871
A:Experimental source: cultivar Columbia
C:Genetics:
A:Map position: 4
A:Introns: 47/3; 201/2; 243/1; 259/2
A:Note: A_TM018A10.10
C:Superfamily: Arabidopsis thaliana hypothetical protein A_TM018A10.10

Query Match 90.5%; Score 19; DB 2; Length 357;
Best Local Similarity 66.7%; Pred. No. 7.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 91 EAGSSS 96

RESULT 29

AB3177
probable N-acetylglucosamine-6-phosphate deacetylase PA3758 [imported] - Pseudomonas
C:Species: Pseudomonas aeruginosa
C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
C:Accession: AB3177
R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.;
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; L
; Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000

A;Title: Complete genome sequence of *Pseudomonas aeruginosa* PA01, an opportunistic pathogen
A;Reference number: A82950; MUID:20437337

A;Accession: A83177

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-363 <STO>

A;Cross-references: GB:AF004794; GB:AE004091; NID:g9949917; PIDN:AG07145.1; GSPDB:GN001

A;Experimental source: strain PA01

C;Genetics:

A;Gene: PA3758

Query Match 90.5%; Score 19; DB 2; Length 363;
Best Local Similarity 66.7%; Pred. No. 7.8e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

||| |

Db 196 EAGASS 201

RESULT 30

T47240

amino acid transport protein arg-1, mitochondrial [imported] - *Neurospora crassa*

C;Species: *Neurospora crassa*

C;Date: 20-Apr-2000 #sequence_revision 20-Apr-2000 #text_change 21-Jul-2000

C;Accession: T47240

R;Liu, Q.; Dunlap, J.C.

Genetics 143, 1163-1174, 1996

A;Title: Isolation and analysis of the arg-13 gene of *Neurospora crassa*.

A;Reference number: 22416; MUID:96400914

A;Accession: T47240

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-363 <LNU>

A;Cross-references: EMBL:L36378; NID:g773383; PIDN:AAC37500.1; PID:g773384

A;Experimental source: strain bda; isolate 30-1

C;Genetics:

A;Gene: arg-13

A;Map position: V

A;Introns: 50/3

C;Keywords: amino acid transport; mitochondrion

Query Match 90.5%; Score 19; DB 2; Length 363;
Best Local Similarity 66.7%; Pred. No. 7.8e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

||| |

Db 14 EAGAAS 19

RESULT 31

E83800

hypothetical protein BH1205 [imported] - *Bacillus halodurans* (strain C-125)

C;Species: *Bacillus halodurans*

C;Date: 01-Dec-2000 #sequence_revision 01-Dec-2000 #text_change 15-Jun-2001

C;Accession: E83800

R;Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; Hira

Nucleic Acids Res. 28, 4317-4331, 2000

A;Title: Complete genome sequence of the alkaliphilic bacterium *Bacillus halodurans* and

A;Reference number: A83650; MUID:20512582; PMID:11058132

A;Accession: E83800

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-365 <STO>

A;Cross-references: GB:AF001511; GB:BA000004; NID:g10173727; PIDN:BA04924.1; GSPDB:GN00

A;Experimental source: strain C-125

C;Genetics:

A;Gene: BH1205

Query Match 90.5%; Score 19; DB 2; Length 365;
Best Local Similarity 66.7%; Pred. No. 7.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

||| |

Db 247 EAGTAS 252

RESULT 32

I40226

3-isopropylmalate dehydrogenase (EC 1.1.1.85) - *Bacillus megaterium*

C;Species: *Bacillus megaterium*

C;Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999

C;Accession: I40226; S38506

R;Meinhardt, F.; Wittchen, K.D.; Buakamp, M.

Appl. Microbiol. Biotechnol. 41, 344-351, 1994

A;Title: Cloning and sequencing of the *lenC* and *npm* genes and a putative *spoIV* gene f

A;Reference number: I40226; MUID:94288995

A;Accession: I40226

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-367 <RES>

A;Cross-references: EMBL:X65184; NID:g414096; PIDN:CAA46295.1; PID:g414097

A;Experimental source: DSM 319

C;Genetics:

A;Gene: *leuc*

C;Function:

A;Pathway: leucine biosynthesis

C;Superfamily: 3-isopropylmalate dehydrogenase

C;Keywords: leucine biosynthesis; NAD; oxidoreductase

Query Match 90.5%; Score 19; DB 1; Length 367;
Best Local Similarity 66.7%; Pred. No. 7.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

||| |

Db 48 EAGSSS 53

RESULT 33

T25452

hypothetical protein B0412.1 - *Caenorhabditis elegans*

C;Species: *Caenorhabditis elegans*

C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999

C;Accession: T25452

R;Bentley, D.

submitted to the EMBL Data Library, December 1996

A;Description: The sequence of *C. elegans* cosmid B0412.

A;Reference number: Z20037

A;Accession: T25452

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-387 <BSN>

A;Cross-references: EMBL:U80953; PIDN:AAB52555.1; GSPDB:GN00021; CESP:B0412.1

A;Experimental source: strain Bristol N2; clone B0412

C;Genetics:

A;Gene: CESP:B0412.1

A;Map position: 3

A;Introns: 110/3; 146/2; 175/1; 213/2; 253/3; 318/1

Query Match 90.5%; Score 19; DB 2; Length 387;
Best Local Similarity 66.7%; Pred. No. 8.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6

||| |

Db 276 EAGSSS 281

RESULT 34

A39429
CAMP response element-binding protein ATF2 - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999
C:Accession: A39429
R:Kageyama, R.; Sasai, Y.; Nakanishi, S.
J. Biol. Chem. 266, 15525-15531, 1991
A:Title: Molecular characterization of transcription factors that bind to the CAMP response element
A:Reference number: A39429; MUID:91332085
A:Accession: A39429
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-389 <KAG>
A:Cross-references: GB:M65148; NID:g206569; PIDN:AAA42013.1; PID:g206570
C:Superfamily: CAMP response element-binding protein 1; fos/jun DNA-binding domain homolog
C:Keywords: DNA binding; nucleus; transcription regulation
F:231-271/Domain: fos/jun DNA-binding domain homology <FJD>

Query Match 90.5%; Score 19; DB 1; Length 389;
Best Local Similarity 66.7%; Pred. No. 8.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |

Db 354 EAGATS 359

RESULT 35

JT0334
acid proteinase (EC 3.4.23.-) PEPI precursor - yeast (Saccharomycopsis fibuligera)
C:Species: Saccharomycopsis fibuligera
C:Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Sep-1999
C:Accession: JT0334
R:Hirata, D.; Fukui, S.; Yamashita, I.
Agric. Biol. Chem. 52, 2647-2649, 1988
A:Title: Nucleotide sequence of the secreted acid protease gene PEPI in the yeast sacc

A:Reference number: JT0334
A:Accession: JT0334
A:Molecule type: DNA
A:Residues: 1-390 <HIR>
C:Genetics:
A:Gene: PEPI
C:Superfamily: pepsin
C:Keywords: aspartic proteinase; hydrolase
F:1-20/Domain: signal sequence #status predicted <SIG>
F:21-390/Product: acid proteinase #status predicted <MAT>
F:93,282/Active site: Asp #status predicted

Query Match 90.5%; Score 19; DB 2; Length 390;
Best Local Similarity 66.7%; Pred. No. 8.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |

Db 266 EAGSSS 271

RESULT 36

F72068
dihydrolipoamide succinyltransferase - Chlamydomophila pneumoniae (strain CWL029)
C:Species: Chlamydomophila pneumoniae, Chlamydia pneumoniae
C:Date: 23-Apr-1999 #sequence_revision 23-Apr-1999 #text_change 05-May-2000
C:Accession: F72068
R:Kallman, S.; Mitchell, W.; Marathe, R.; Lammel, C.; Fan, J.; Olinger, L.; Grimwood, J.; Nature Genet. 21, 385-389, 1999
A:Title: Comparative genomes of Chlamydia pneumoniae and C. trachomatis.
A:Reference number: A72000; MUID:99206606
A:Accession: F72068
A:Status: preliminary
A:Molecule type: DNA

A:Residues: 1-393 <ARN>

A:Cross-references: GB:AE001637; GB:AE001363; NID:g4376807; PIDN:AAI8667.1; PID:g437
A:Experimental source: strain CWL029

C:Genetics:

A:Gene: sucB_2

C:Superfamily: dihydrolipoamide acetyltransferase; lipoyl/biotin-binding homology

Query Match 90.5%; Score 19; DB 2; Length 393;
Best Local Similarity 66.7%; Pred. No. 8.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |

Db 101 EAGSSS 106

RESULT 37

C86556
dihydrolipoamide succinyltransferase [imported] - Chlamydomophila pneumoniae (strain J1
C:Species: Chlamydomophila pneumoniae, Chlamydia pneumoniae
C:Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 23-Mar-2001
C:Accession: C86556
R:Shirai, M.; Hirakawa, H.; Kimoto, M.; Tabuchi, M.; Kishi, F.; Ouchi, K.; Shiba, T.; Nucleic Acids Res. 28, 2311-2314, 2000
A:Title: Comparison of whole genome sequences of chlamydia pneumoniae J138.
A:Reference number: A86491; MUID:20330349
A:Accession: C86556
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-393 <STO>
A:Cross-references: GB:BA000008; NID:g8978898; PIDN:BA098733.1; GSPDB:GN00142
A:Experimental source: strain J138
C:Genetics:
A:Gene: sucB_2

C:Superfamily: dihydrolipoamide acetyltransferase; lipoyl/biotin-binding homology

Query Match 90.5%; Score 19; DB 2; Length 393;
Best Local Similarity 66.7%; Pred. No. 8.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |

Db 101 EAGSSS 106

RESULT 38

AD1559
glycine betaine ABC transporter (ATP-binding protein) homolog gbuA [imported] - Liste
C:Species: Listeria innocua
C:Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 14-Dec-2001
C:Accession: AD1559
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloec
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi,
D.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapat, G.; Madueno, E.; Maitournam, A.;
ok, C.; Schlueter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wenla
A:Title: Comparative genomics of Listeria species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669
A:Accession: AD1559
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-397 <GLA>

A:Cross-references: GB:AL592022; PIDN:CA096244.1; PID:g16413472; GSPDB:GN00178

A:Experimental source: strain Clip11262

C:Genetics:

A:Gene: gbuA

C:Superfamily: glycine betaine/proline transport protein prov; ATP-binding cassette h

Query Match 90.5%; Score 19; DB 2; Length 397;

```
Best Local Similarity 66.7%; Pred. No. 8.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
   ||| |
Db 301 EAGTSS 306

RESULT 39
AF1201
glycine betaine ABC transporter (ATP-binding protein) homolog gbaA [imported] - Listeria
C:Species: Listeria monocytogenes
C:Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 14-Dec-2001
C:Accession: AF1201
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloeker,
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi, H.
D.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A:Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.; Maitournam, A.; Ma
ok, C.; Schluter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehland,
A:Title: Comparative genomics of Listeria species.
A:Reference number: AB1077; MUID:21537279; PMID:11679669
A:Accession: AF1201
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-397 <GLA>
A:Cross-references: GB:NC_003210; PIDN:CAC99092.1; PID:g16410416; GSPDB:GN00177
A:Experimental source: strain EGD-e
C:Genetics:
A:Gene: gbaA
C:Superfamily: glycine betaine/proline transport protein prov; ATP-binding cassette hom

Query Match 90.5%; Score 19; DB 2; Length 397;
Best Local Similarity 66.7%; Pred. No. 8.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
   ||| |
Db 301 EAGTSS 306

RESULT 40
B72778
probable processing proteinase APE0212 - Aeropyrum pernix (strain K1)
C:Species: Aeropyrum pernix
C:Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Jun-2000
C:Accession: B72778
R:Kawarabayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takah
awa, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, T.; Kudo, Y.; Yamazaki, J.; K
DNA Res. 6, 83-101, 1999
A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyr
A:Reference number: A72450; MUID:99310339
A:Accession: B72778
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-403 <RAW>
A:Cross-references: DDBJ:AP000058; NID:G5103380; PIDN:BAA79124.1; PID:g5103603
A:Experimental source: strain K1
C:Genetics:
A:Gene: APE0212
C:Superfamily: mitochondrial processing peptidase alpha chain

Query Match 90.5%; Score 19; DB 2; Length 403;
Best Local Similarity 66.7%; Pred. No. 8.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
   ||| |
Db 266 EAGATS 271
```

```
RESULT 41
A54813
CAMP receptor CAR4 - slime mold (Dictyostelium discoideum)
C:Species: Dictyostelium discoideum
C:Date: 23-Mar-1995 #sequence_revision 05-Apr-1995 #text_change 07-May-1999
C:Accession: A54813
R:Louis, J.M.; Ginsburg, G.T.; Kimmel, A.R.
Genes Dev. 8, 2086-2096, 1994
A:Title: The CAMP receptor CAR4 regulates axial patterning and cellular differentiati
A:Reference number: A54813; MUID:95047357
A:Accession: A54813
A:Status: preliminary; nucleic acid sequence not shown; not compared with conceptual
A:Molecule type: DNA
A:Residues: 1-443 <LOU>
C:Genetics:
A:Gene: CAR4
C:Keywords: CAMP binding

Query Match 90.5%; Score 19; DB 2; Length 443;
Best Local Similarity 66.7%; Pred. No. 9.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
   ||| |
Db 72 EAGSTS 77

RESULT 42
S35783
glycoprotein gX - bovine herpesvirus 1
C:Species: bovine herpesvirus 1
C:Date: 09-Jun-1994 #sequence_revision 12-May-1995 #text_change 08-Oct-1999
C:Accession: S35783
R:Audonnet, J.
submitted to the EMBL Data Library, June 1993
A:Reference number: S35782
A:Accession: S35783
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-444 <AUD>
A:Cross-references: EMBL:Z23068; NID:g312185; PIDN:CAA80603.1; PID:g312187
C:Superfamily: pseudorabies virus glycoprotein gX
C:Keywords: glycoprotein

Query Match 90.5%; Score 19; DB 2; Length 444;
Best Local Similarity 66.7%; Pred. No. 9.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
   ||| |
Db 267 EAGSAS 272

RESULT 43
T02804
hypothetical protein L2602.2 [imported] - Leishmania major (strain Friedlin)
C:Species: Leishmania major
C:Date: 24-Mar-1999 #sequence_revision 24-Mar-1999 #text_change 19-May-2000
C:Accession: H81456; T02804
R:Myler, P.J.; Audleman, L.; deVos, T.; Hixson, G.; Kiser, P.; Lemley, C.; Magness, C
Proc. Natl. Acad. Sci. U.S.A. 96, 2902-2906, 1999
A:Title: Leishmania major Friedlin chromosome 1 has an unusual distribution of protei
A:Reference number: A81455; MUID:99178987
A:Accession: H81456
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-458 <PYL>
A:Cross-references: GB:AE001274; NID:g3264850; PIDN:AAC24627.1; PID:g2995580; GSPDB:G
A:Experimental source: strain MHOW/IL/81/Friedlin
C:Genetics:
A:Gene: L2602.2
```


A: Map position: 1

C: Superfamily: Leishmania major hypothetical protein L2602.2

Query Match 90.5%; Score 19; DB 2; Length 458;
Best Local Similarity 66.7%; Pred. No. 9.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
DB 10 EAGTAS 15

RESULT 44

D85438
hypothetical protein At4g37110 [imported] - Arabidopsis thaliana

C: Species: Arabidopsis thaliana (mouse-ear cress)

C: Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 16-Feb-2001

C: Accession: D85438

R: anonymous, The European Union Arabidopsis Genome Sequencing Consortium, The Cold Spring Nature 402, 769-777, 1999

A: Title: Sequence and analysis of chromosome 4 of the plant Arabidopsis thaliana.

A: Reference number: A85001; MUID: 20083488

A: Accession: D85438

A: Status: preliminary

A: Molecule type: DNA

A: Residues: 1-462 <STO>

A: Cross-references: GB:NC_001268; NID: g7270660; PIDN: CAB80377.1; GSPDB: GN00140

C: Genetics:

A: Gene: At4g37110

A: Map position: 4

Query Match

Best Local Similarity 90.5%; Score 19; DB 2; Length 462;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
DB 108 EAGAAS 113

RESULT 45

C87629

major facilitator family transporter CC3069 [imported] - Caulobacter crescentus

C: Species: Caulobacter crescentus

C: Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 10-May-2001

C: Accession: C87629

R: Nierman, W.C.; Feidblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J. B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kolon, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.M. Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001

A: Title: Complete Genome Sequence of Caulobacter crescentus.

A: Reference number: A87249; MUID: 21173698; PMID: 11259647

A: Accession: C87629

A: Status: preliminary

A: Molecule type: DNA

A: Residues: 1-469 <STO>

A: Cross-references: GB:AE005673; NID: g13424719; PIDN: AAK25031.1; GSPDB: GN00148

C: Genetics:

A: Gene: CC3069

C: Superfamily: lincomycin-resistance protein lmrB

Query Match

Best Local Similarity 90.5%; Score 19; DB 2; Length 469;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
DB 394 EAGAAS 399

RESULT 46

D70853

hypothetical protein Rv3088 - Mycobacterium tuberculosis (strain H37RV)

C: Species: Mycobacterium tuberculosis

C: Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999

C: Accession: D70853

R: Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon ; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S. Nature 393, 537-544, 1998

A: Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.

A: Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno

A: Reference number: A70500; MUID: 98295987

A: Accession: D70853

A: Status: preliminary; nucleic acid sequence not shown; translation not shown

A: Molecule type: DNA

A: Residues: 1-474 <COL>

A: Cross-references: GB:AL021309; GB:AL123456; NID: g3261510; PIDN: CAA16146.1; PID: e124

A: Experimental source: strain H37RV

C: Genetics:

A: Gene: Rv3088

Query Match

Best Local Similarity 90.5%; Score 19; DB 2; Length 474;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
DB 460 EAGTTS 465

RESULT 47

S08384

modulation protein nodT - Rhizobium leguminosarum plasmid pIJ1089

C: Species: Rhizobium leguminosarum

C: Date: 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change 18-Sep-1998

C: Accession: S08384

R: Economou, A.; Hamilton, W.D.O.; Johnston, A.W.B.; Downie, J.A. EMBO J. 9, 349-354, 1990

A: Title: The Rhizobium modulation gene nodO encodes a Ca²⁺-binding protein that is ex

A: Reference number: S08384; MUID: 90151607

A: Accession: S08384

A: Status: preliminary; nucleic acid sequence not shown; translation not shown

A: Molecule type: DNA

A: Residues: 1-482 <ECO>

A: Cross-references: EMBL: X17285

A: Note: the nucleotide sequence was submitted to the EMBL Data Library, December 1989

C: Genetics:

A: Genome: plasmid pIJ1089

C: Superfamily: modulation protein nodT

Query Match

Best Local Similarity 90.5%; Score 19; DB 2; Length 482;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
DB 201 EAGAAS 206

RESULT 48

S10133

modulation protein nodT - Rhizobium leguminosarum bv. viciae

C: Species: Rhizobium leguminosarum bv. viciae

C: Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 29-Sep-1999

C: Accession: S10133

R: Surin, B.P.; Watson, J.M.; Hamilton, W.D.O.; Economou, A.; Downie, J.A. Mol. Microbiol. 4, 245-252, 1990

A: Title: Molecular characterization of the modulation gene, nodT, from two biovars of

A: Reference number: S08616; MUID: 90251164

A: Accession: S10133

A:Molecule type: DNA
A:Residues: 1-482 <SUR>
A:Cross-references: EMBL:X17285; NID:g46251; PIDN:CAA35177.1; PID:g581512
C:Genetics:
A:Gene: nodT
A:Start codon: TTG
C:Superfamily: nodulation protein nodT

Query Match 90.5%; Score 19; DB 2; Length 482;
Best Local Similarity 66.7%; Pred. No. 1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 201 EAGAS 206

RESULT 49

T41039
probable transcription initiation factor Iif, alpha subunit - fission yeast (Schizosaccharomyces pombe)
C:Species: Schizosaccharomyces pombe
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 07-Dec-1999
C:Accession: T41039
R:Lyne, M.; Rajandream, M.A.; Barrell, B.G.; Hilbert, H.; Duesterhoeft, A.
submitted to the EMBL Data Library, December 1998
A:Reference number: Z21966
A:Accession: T41039
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-490 <LYN>
A:Cross-references: EMBL:AL034491; PIDN:CAA22493.1; GSPDB:GN00068; SPDB:SPCC1620.09c
A:Experimental source: strain 972h-; cosmid c1620
C:Genetics:
A:Gene: SPDB:SPCC1620.09c
A:Map position: 3
A:Introns: 180/3; 212/3
C:Keywords: transcription initiation

Query Match 90.5%; Score 19; DB 2; Length 490;
Best Local Similarity 66.7%; Pred. No. 1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 234 EAGAS 239

RESULT 50

S54536
probable membrane protein YDR240c - yeast (Saccharomyces cerevisiae)
N:Alternate names: hypothetical protein YD8419.07c
C:Species: Saccharomyces cerevisiae
C:Date: 08-Jul-1995 #sequence_revision 01-Sep-1995 #text_change 04-Mar-2000
C:Accession: S54536
R:Oliver, K.; Harris, D.
submitted to the EMBL Data Library, May 1995
A:Reference number: S54530
A:Accession: S54536
A:Molecule type: DNA
A:Residues: 1-492 <OLI>
A:Cross-references: EMBL:Z49701; NID:g817819; PID:g817826; GSPDB:GN00004; MIPS:YDR240c
A:Experimental source: strain AB972
C:Genetics:
A:Gene: MIPS:YDR240c
A:Map position: 4R
C:Superfamily: Saccharomyces cerevisiae probable membrane protein YDR240c
C:Keywords: transmembrane protein
F:119-135/Domain: transmembrane #status predicted <TMM>

Query Match 90.5%; Score 19; DB 2; Length 492;

Best Local Similarity 66.7%; Pred. No. 1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 325 EAGATS 330

Search completed: August 30, 2002, 15:09:49
Job time: 585 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: August 30, 2002, 15:06:44 ; Search time 13.51 Seconds
(without alignments)
17.196 Million cell updates/sec

Title: BASK-853-CLAIM4

Perfect score: 21

Sequence: 1 eaggxs 6

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 105224 seqs, 38719550 residues

Total number of hits satisfying chosen parameters: 105224

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 50 summaries

Database : SwissProt_40.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19	90.5	303	1 HISL_HAEIN	P43953 haemophilus
2	19	90.5	322	1 YK02 YEAST	P36118 saccharomyc
3	19	90.5	323	1 MAFL MOUSE	P54841 mus musculus
4	19	90.5	323	1 MAFL RAT	P54842 rattus norv
5	19	90.5	331	1 MACS HUMAN	P29966 homo sapien
6	19	90.5	352	1 NPIL ASPOR	P46076 aspergillus
7	19	90.5	363	1 AR13 NEUCR	Q01356 neurospora
8	19	90.5	367	1 LEU3 BACME	P41019 bacillus me
9	19	90.5	390	1 CARP_SACFI	P22929 saccharomyc
10	19	90.5	443	1 CAR4_DICDI	Q9TX43 dictyosteli
11	19	90.5	444	1 VGLX HSVBS	Q08103 bovine herp
12	19	90.5	474	1 YU88 MYCTU	O53305 mycobacteri
13	19	90.5	482	1 NODT_RHLV	P15727 rhizobium 1
14	19	90.5	487	1 ATF2_RAT	Q00969 rattus norv
15	19	90.5	561	1 MERA_ACICA	Q52109 acinetobact
16	19	90.5	561	1 MERA_ENTAG	P94702 enterobacte
17	19	90.5	578	1 AC22_SYRCO	P46105 streptomyce
18	19	90.5	636	1 DNK2_SYNY3	P22358 synecocyst
19	19	90.5	638	1 TOXA_PSEAE	P11439 pseudomonas
20	19	90.5	643	1 SGT1_ARATH	Q915m5 arabidopsis
21	19	90.5	692	1 EOMD_XENLA	P79944 xenopus lae
22	19	90.5	883	1 PGCB_MOUSE	Q61361 mus musculus
23	19	90.5	883	1 PGCB_RAT	P55068 rattus norv
24	19	90.5	908	1 SRCA_RABIT	P22358 mycobacteri
25	19	90.5	1012	1 POLS_IBDVA	P08364 avian infec
26	19	90.5	1021	1 TSCC_HUMAN	P55017 homo sapien
27	19	90.5	1027	1 ISWI_DROME	Q24368 drosophila
28	19	90.5	1058	1 PMA1_DICDI	P54679 dictyosteli
29	19	90.5	1069	1 CNA_ARATH	Q9SZR1 arabidopsis
30	19	90.5	1183	1 ACNA_ARATH	Q53654 staphylococ
31	19	90.5	1237	1 B3A2_MOUSE	P13808 mus musculus
32	19	90.5	1262	1 MYO6_HUMAN	Q9um54 homo sapien
33	19	90.5	1265	1 DYNA_DROME	P13496 drosophila

ALIGNMENTS

RESULT 1

HISL_HAEIN

ID HISL_HAEIN

AC P43853;

DT 01-NOV-1995 (Rel. 32, Created)

DT 01-NOV-1995 (Rel. 32, Last sequence update)

DT 16-OCT-2001 (Rel. 40, Last annotation update)

DE ATP phosphoribosyltransferase (EC 2.4.2.17).

GN HISG OR HI0468.

OS Haemophilus influenzae.

OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;

OC Haemophilus.

OX NCBI_TaxID=727;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=RD / KW20 / ATCC 51907;

RX MEDLINE=95350630; PubMed=7542800;

RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,

RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,

RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,

RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,

RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,

RA Utterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,

RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,

RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,

RA Venter J.C.;

RT "Whole-genome random sequencing and assembly of Haemophilus

RT influenzae Rd.";

RL Science 269:496-512(1995).

CC -1- CATALYTIC ACTIVITY: 1-(5-phospho-D-ribose)-ATP + diphosphate =

CC -1- ATP + 5-phospho-alpha-D-ribose 1-diphosphate.

CC -1- PATHWAY: FIRST STEP IN HISTIDINE BIOSYNTHETIC PATHWAY. IS VERY

CC -1- IMPORTANT IN THE REGULATION OF HISTIDINE METABOLISM.

CC -1- SUBUNIT: HOMOHXAMER (BY SIMILARITY).

CC -1- SUBCELLULAR LOCATION: Cytoplasmic (BY SIMILARITY).

CC -1- SIMILARITY: BELONGS TO THE ATP PHOSPHORIBOSYLTRANSFERASE FAMILY.

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CC -----

CC EMBL; U32729; AAC2127.1; -.

CC TIGR; HI0468; -.

CC InterPro; IPR001348; HisG.

CC Pfam; PF01634; HisG; 1.

CC PROSITE; PD003516; HisG; 1.

CC PROSITE; PS01316; ATP_P_PHORIBOSYLTR; 1.

Q64331 mus musculus
Q92628 homo sapien
P53564 mus musculus
P22523 escherichia
P39880 homo sapien
Q09332 drosophila
P48633 yersinia en
P81671 pinus pinas
P00294 capsella bu
O14904 homo sapien
P35137 bacillus su
Q08183 bos taurus
Q07199 mycobacteri
Q92qk5 rhizobium m
P58233 escherichia
O51466 pseudomonas
P34469 caenorhabdi

1 MYO6_MOUSE
1 Y232_HUMAN
1 CUT1_MOUSE
1 MUKB_ECOLI
1 CUT3_HUMAN
1 UGGG_DROME
1 HMP2_YEREN
1 CLPA_PINPS
1 PLAS_CAPBU
1 WN14_HUMAN
1 PP1B_BACSU
1 RI57_BOVIN
1 DUT_MYCTU
1 MOAE_RHIME
1 CEST_ECO57
1 FLIN_PSEAE
1 YMH2_CAEEL

1265
1278
1395
1486
1505
1548
2035
30
99
123
143
147
154
155
156
157
159

19
19
19
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19
18
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18
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90.5
90.5
90.5
90.5
90.5
90.5
90.5
85.7
85.7
85.7
85.7
85.7
85.7
85.7
85.7
85.7

KW Histidine biosynthesis; Transferase; Glycosyltransferase;
 KW Complete proteome.
 SQ SEQUENCE 303 AA; 33821 MW; 08C14D1F6E98A31D CRC64;

Query Match 90.5%; Score 19; DB 1; Length 303;
 Best Local Similarity 66.7%; Pred. No. 2.5e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 |||||
 Db 287 EAGASS 292

RESULT 2

YK02_YEAST STANDARD; PRT; 322 AA.
 AC P36118;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DE Hypothetical 36.6 kDa protein in YPT52-DBP7 intergenic region.
 GN YK022C.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 OX NCBI_TaxID=4932;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C;
 RL Submitted (MAR-1994) to the EMBL/GenBank/DBJ databases.
 CC -----
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 CC -----

EMBL; Z28247; GAA82094.1; -
 DR PIR; S38091;
 DR SGD; S0001730; YK022C.
 KW Hypothetical protein.
 SQ SEQUENCE 322 AA; 36647 MW; D7A601A46839244C CRC64;

Query Match 90.5%; Score 19; DB 1; Length 322;
 Best Local Similarity 66.7%; Pred. No. 2.6e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 |||||
 Db 119 EAGSSS 124

RESULT 3

MAF1_MOUSE STANDARD; PRT; 323 AA.
 AC P54841;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Transcription factor MAF1 (Segmentation protein KR) (Kreisler).
 GN MAFB OR MAF1 OR KRML.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC MEDLINE=95094266; PubMed=8001130;

RA Cordes S.P., Barsh G.S.;
 RT "The mouse segmentation gene *kr* encodes a novel basic domain-leucine
 RL zipper transcription factor.";
 RL Cell 79:1025-1034(1994).
 CC -!- FUNCTION: MAY PLAY AN EARLY ROLE IN AXIAL PATTERNING (HINDBRAIN
 CC SEGMENTATION).
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- TISSUE SPECIFICITY: MOST ABUNDANT IN KIDNEY, GUT, LUNG, AND BRAIN.
 CC -!- DEVELOPMENTAL STAGE: DETECTABLE AT 8.0 DPC (ONE SOMITE) AS A BAND
 CC IN THE CAUDAL HINDBRAIN, AND BY 8.5 DPC (SIX TO EIGHT SOMITES),
 CC THE HIGH LEVEL DOMAIN EXHIBITS A SHARP ROSTRAL EDGE COINCIDENT
 CC WITH THE R4/R5 BOUNDARY AND A DIFFUSE CAUDAL EDGE LOCATED MIDWAY
 CC THROUGH R6.
 CC -!- SIMILARITY: BELONGS TO THE BZIP FAMILY. MAF SUBFAMILY.
 CC -----
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 CC -----

EMBL; L36435; AAA65689.1; -

DR HSSP; P05412; LJUN.
 DR TRANSFAC; T01439; -
 DR MGD; MGI:104555; MafB.
 DR InterPro; IPR001871; bZIP.
 DR Pfam; PF03131; bZIP_Maf; 1.
 DR SMART; SM00338; BRL2; 1.
 KW Transcription regulation; DNA-binding; Nuclear protein.
 FT DOMAIN 131 143 POLY-HIS.
 FT DOMAIN 158 167 POLY-HIS.
 FT DNA_BIND 238 264 BASIC MOTIF.
 FT DOMAIN 266 287 LEUCINE-ZIPPER.
 FT MUTAGEN 248 248 N->S: LOSS OF TRANSCRIPTIONAL ACTIVITY.
 SQ SEQUENCE 323 AA; 35809 MW; D77AE07ABD9C2AD2 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 323;

Best Local Similarity 66.7%; Pred. No. 2.6e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 |||||
 Db 308 EAGSTS 313

RESULT 4

MAF1_RAT STANDARD; PRT; 323 AA.
 AC P54842;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Transcription factor MAF1.
 GN MAFB OR MAF1.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Wistar; Tissue=Liver;
 RX MEDLINE=97190228; PubMed=9038383;
 RA Sakai M., Imaki J., Yoshida K., Ogata A., Matsushima-Hibaya Y.,
 RA Kuboki Y., Nishizawa M., Nishi S.;
 RT "Rat *maf* related genes: specific expression in chondrocytes, lens and
 RL spinal cord.";
 RL Oncogene 14:745-750(1997).
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- SIMILARITY: BELONGS TO THE BZIP FAMILY. MAF SUBFAMILY.
 CC -----

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DR EMBL; U56241; AAB50062.1; -
DR HSSP; P05412; LJUN.
DR InterPro; IPR001871; bZIP.
DR Pfam; PF03131; bZIP_Maf; 1.
DR SMART; SM00338; BRLZ; 1.
KW Transcription regulation; DNA-binding; Nuclear protein.
FT DOMAIN 131 143 POLY-HIS.
FT DOMAIN 158 167 POLY-HIS.
FT DOMAIN 194 201 POLY-ALA.
FT DNA_BIND 238 264 BASIC MOTIF.
FT DOMAIN 266 287 LEUCINE-ZIPPER.
SQ SEQUENCE 323 AA; 35792 MW; 6E386340D1F840A5 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 323;
Best Local Similarity 66.7%; Pred. No. 2.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 308 EAGSTS 313

RESULT 5
ID MACS_HUMAN STANDARD; PRT; 331 AA.
AC P29966;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-JUL-1993 (Rel. 26, Last sequence update)
DE Myristoylated alanine-rich C-kinase substrate (MARCKS) (Protein kinase
DE C substrate, 80 kDa protein, light chain) (PKCSL) (80K-L protein).
GN MACS.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE-91317795; PubMed-1860846;
RA Harlan D.M., Graff J.M., Stumpo D.J., Eddy R.L. Jr., Shows T.B.,
RA Boyie J.M., Blackshear P.J.;
RT "The human myristoylated alanine-rich C kinase substrate (MARCKS)
RT gene (MACS). Analysis of its gene product, promoter, and chromosomal
RT localization.";
RL J. Biol. Chem. 266:14399-14405(1991).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE-93052291; PubMed-1427823;
RA Sakai K., Hirai M., Kudo J., Minoshima S., Shimizu N.;
RT "Molecular cloning and chromosomal mapping of a cDNA encoding human
RT 80K-L protein: major substrate for protein kinase C.";
RL Genomics 14:175-178(1992).
CC -!- FUNCTION: MARCKS IS THE MOST PROMINENT CELLULAR SUBSTRATE FOR
CC PROTEIN KINASE C. THIS PROTEIN BINDS CALMODULIN, ACTIN, AND
CC SYNAPSIN. MARCKS IS A FILAMENTOUS (F) ACTIN CROSS-LINKING PROTEIN.
CC -!- PTM: PHOSPHORYLATION BY PKC REPLACES MARCKS FROM THE MEMBRANE. IT
CC ALSO INHIBITS THE F-ACTIN CROSS-LINKING ACTIVITY.
CC -!- SIMILARITY: BELONGS TO THE MARCKS FAMILY.

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DR EMBL; M68956; AAA59555.1; -
DR EMBL; M68955; AAA59554.1; -
DR EMBL; D10522; BAA01392.1; -
DR PIR; A38873; A38873.
DR MIM; 177061; -
DR InterPro; IPR002101; MARCKS.
DR Pfam; PF02063; MARCKS; 1.
DR PRINTS; PR00963; MARCKS.
DR PROSITE; PS00826; MARCKS_1; 1.
DR PROSITE; PS00827; MARCKS_2; 1.
KW Phosphorylation; Myristate; Calmodulin-binding; Actin-binding;
KW Membrane.
FT INIT_MET 0 0 BY SIMILARITY.
FT LIPID 1 1 MYRISTATE (BY SIMILARITY).
FT DOMAIN 151 175 CALMODULIN-BINDING (PSD).
FT MOD_RES 158 162 PHOSPHORYLATION (BY PKC).
FT MOD_RES 162 162 PHOSPHORYLATION (BY PKC).
FT MOD_RES 166 166 PHOSPHORYLATION (BY PKC).
FT MOD_RES 169 169 PHOSPHORYLATION (BY PKC).
FT CONFLICT 83 83 S -> A (IN REF. 1).
FT CONFLICT 118 118 A -> P (IN REF. 1).
FT CONFLICT 233 233 P -> S (IN REF. 1).
FT CONFLICT 286 307 LVCPRRGSGPRGGRRSLNQ (IN REF. 1).
SQ SEQUENCE 331 AA; 31413 MW; BCC837D586581774 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 331;
Best Local Similarity 66.7%; Pred. No. 2.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 205 EGAAS 210

RESULT 6
ID NPIL_ASPOR STANDARD; PRT; 352 AA.
AC P46076;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-FEB-1996 (Rel. 33, Last annotation update)
DE Neutral protease II precursor (EC 3.4.24.39) (Deuterolysin) (NPIL).
OS Aspergillus oryzae.
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Eurotiomycetes;
OC Eurotiales; Trichocomaceae; mitosporic Trichocomaceae; Aspergillus.
OX NCBI_TaxID=5062;
RN [1]
RP SEQUENCE FROM N.A., AND SEQUENCE OF 176-210; 279-281 AND 304-341.
RC STRAIN-ATCC 20386;
RX MEDLINE-91360097; PubMed-1886621;
RA Tatsumi H., Murakami S., Tsuji R.F., Ishida Y., Murakami K.,
RA Masaki A., Kawabe H., Arimura H., Nakano E., Motai H.;
RT "Cloning and expression in yeast of a cDNA clone encoding Aspergillus
RT oryzae neutral protease II, a unique metalloprotease.";
RL Mol. Gen. Genet. 228:97-103(1991).
CC -!- FUNCTION: THERMOSTABLE METALLOPROTEASE. SHOWS HIGH ACTIVITIES ON
CC BASIC NUCLEAR SUBSTRATES SUCH AS HISTONE AND PROTAMINE.
CC -!- CATALYTIC ACTIVITY: Preferential cleavage of bonds with
CC hydrophobic residues in p1'; also 3-Asn-1-Gln-4 and 8-Gln-1-Ser-9
CC bonds in insulin B chain.
CC -!- COFACTOR: BINDS 1 ZINC ION.
CC -!- PTM: PROBABLY POSSESSES THREE DISULFIDE BONDS.
CC -!- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M35 (ZINC
CC METALLOPROTEASE); ALSO KNOWN AS THE DEUTEROLYSIN SUBFAMILY.

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DR EMBL; S53810; AAB19701.1; -.
DR MEROPS; M35.002; -.
DR InterPro; IPR001384; Peptidase_M35.
DR InterPro; IPR000130; Zn_MTpeptidse.
DR Pfam; PF02102; Peptidase_M35; 1.
DR PRINTS; PR00768; DEUTEROLYSIN.
DR PROSITE; PS00142; ZINC_PROTEASE; 1.
KW Hydrolase; Metalloprotease; Zinc; Signal; Zymogen.
FT SIGNAL 1 19 POTENTIAL.
FT PROPEP 20 175
FT CHAIN 176 352 NEUTRAL PROTEASE II.
FT METAL 303 303 ZINC (CATALYTIC) (BY SIMILARITY).
FT ACT_SITE 304 304 BY SIMILARITY.
FT METAL 307 307 ZINC (CATALYTIC) (BY SIMILARITY).
SQ SEQUENCE 352 AA; 37517 MW; 070C5131335B7F44 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 352;
Best Local Similarity 66.7%; Pred. No. 2.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 240 EAGSTS 245

RESULT 7
AR13_NEUCR ID ARI3_NEUCR STANDARD; PRT; 363 AA.
AC Q01356;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Amino acid transporter arg-13.
GN ARG-13.
OS Neurospora crassa.
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
OC Sordariales; Sordariaceae; Neurospora.
OX NCBI_TaxID=5141;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BDA;
RX MEDLINE=96400914; PubMed=8807290;
RA Liu Q., Dunlap J.C.;
RT "Isolation and analysis of the arg-13 gene of Neurospora crassa."
RL Genetics 143:1163-1174(1996).
[2]
RP SEQUENCE FROM N.A.
RA Liu Q., Luo X.;
RT "Phenotypic rescue of Saccharomyces cerevisiae arg11 mutant by
RT Neurospora crassa arg-13 cDNA."
RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: REQUIRED FOR ARGinine BIOSYNTHESIS. MAY PARTICIPATE IN
CC THE EXPORT OF MATRIX-MADE ORNITHINE INTO THE CYTOSOL (POTENTIAL).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein. Mitochondrial
CC inner membrane (Potential).
CC -!- DOMAIN: COMPOSED OF THREE HOMOLOGOUS DOMAINS.
CC -!- SIMILARITY: BELONGS TO THE MITOCHONDRIAL CARRIER FAMILY.

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DR EMBL; L36378; AAC37500.1; -.
Query Match 90.5%; Score 19; DB 1; Length 363;
Best Local Similarity 66.7%; Pred. No. 3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

DR EMBL; AF279268; AAF87777.1; -.
DR InterPro; IPR001993; Mitoch_carrier.
DR Pfam; PF00153; mito_carr; 3.
DR PROSITE; PS00215; MITOCH_CARRIER; 2.
KW Mitochondrion; Inner membrane; Repeat; Transmembrane; Transport.
FT TRANSMEM 141 161 POTENTIAL.
FT TRANSMEM 266 283 POTENTIAL.
FT TRANSMEM 334 353 POTENTIAL.
SQ SEQUENCE 363 AA; 39401 MW; 8B87A937F6D37DC0 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 363;
Best Local Similarity 66.7%; Pred. No. 3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 14 EAGAAS 19

RESULT 8
LEU3_BACME ID LEU3_BACME STANDARD; PRT; 367 AA.
AC P41019;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE 3-isopropylmalate dehydrogenase (EC 1.1.1.85) (Beta-IPM dehydrogenase)
DE (IMDH) (3-IPM-DH).
GN LEUB OR LEUC.
OS Bacillus megaterium.
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Bacillus.
OX NCBI_TaxID=1404;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=DSM 319;
RX MEDLINE=94288995; PubMed=7764969;
RA Meinhardt F., Busskamp M., Wittchen K.D.;
RT "Cloning and sequencing of the leu C and npr M genes and a putative
RT spo IV gene from Bacillus megaterium DSM319."
RL Appl. Microbiol. Biotechnol. 41:344-351(1994).
CC -!- CATALYTIC ACTIVITY: 3-CARBOXY-2-HYDROXY-4-METHYLPENTANOATE +
CC NAD(+) = 3-CARBOXY-4-METHYL-2-OXOPENTANOATE + NADH (THE PRODUCT
CC DECARBOXYLATES TO 4-METHYL-2-OXOPENTANOATE).
CC -!- PATHWAY: THIRD STEP IN LEUCINE BIOSYNTHESIS.
CC -!- SUBUNIT: HOMODIMER.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic.
CC -!- SIMILARITY: BELONGS TO THE ISOCITRATE AND ISOPROPYLMALATE
CC DEHYDROGENASES FAMILY.

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DR EMBL; X65184; CAA46295.1; -.
DR PIR; S38506; S38506.
DR HSPSP; P12010; 2AYQ.
DR InterPro; IPR001804; Isodh.
DR Pfam; PF00180; isodh; 1.
DR PROSITE; PS00470; IDH_IMDH; 1.
KW Oxidoreductase; Leucine biosynthesis; NAD.
SQ SEQUENCE 367 AA; 39942 MW; DC04D48E0EEAB0DD CRC64;

Query Match 90.5%; Score 19; DB 1; Length 367;
Best Local Similarity 66.7%; Pred. No. 3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;


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QY 1 eagxxs 6
DB 48 EAGSS 53

RESULT 9
CARP_SACFI STANDARD; PRT; 390 AA.
AC P22929;
DT 01-AUG-1991 (Rel. 19, Created)
DT 01-AUG-1991 (Rel. 19, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Acid protease precursor (EC 3.4.23.-).
GN PEPI.
OS Saccharomycopsis fibuligera (Yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycopsidaceae; Saccharomycopsis.
OX NCBI_TaxID=4944;
RN [1]
SEQUENCE FROM N.A.
RA Hirata D., Fukui S., Yamashita I.;
RT "Nucleotide sequence of the secretible acid protease gene PEPI in the
RL yeast Saccharomycopsis fibuligera.";
RL Agric. Biol. Chem. 52:2647-2649(1988).
CC -! SIMILARITY: BELONGS TO PEPTIDASE FAMILY A1; ALSO KNOWN AS THE
CC EUKARYOTIC ASPARTYL PROTEASES FAMILY.
CC -----
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CC -----
CC EMBL; D00313; BAA00215.1; -
CC PIR; J03334; J03334.
CC HSP; P32329; IYPS.
CC InterPro; IPR001969; Asp_protease.
CC InterPro; IPR001461; Pepsin.
CC Pfam; PF00026; asp; 1.
CC PRINTS; PR00792; PEPSIN.
CC PROSITE; PS00141; ASP_PROTEASE; 2.
KW Hydroxylase; Aspartyl protease; Signal.
FT SIGNAL 1 18 POTENTIAL.
FT CHAIN 19 390 ACID PROTEASE.
FT ACT_SITE 93 93 BY SIMILARITY.
FT ACT_SITE 282 282 BY SIMILARITY.
SQ SEQUENCE 390 AA; 41263 MW; 350BF97116C54796 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 390;
Best Local Similarity 66.7%; Pred. No. 3.2e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
DB 266 EAGSS 271

RESULT 10
CAR4_DICDI STANDARD; PRT; 443 AA.
ID CAR4_DICDI
AC Q9TX43;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DE Cyclic AMP receptor 4.
DE CARD OR CAR4.
OS Dictyostellium discoideum (Slime mold).
OC Eukaryota; Mycetozoa; Dictyostelidida; Dictyostellium.
OX NCBI_TaxID=44689;
RN [1]

SEQUENCE FROM N.A.
STRAIN-NC-4;
RX MEDLINE=95047357; PubMed=7958880;
RA Louis J.M., Ginsburg G.T., Kimmel A.R.;
RT "The CAMP receptor CAR4 regulates axial patterning and cellular
RL differentiation during late development of Dictyostellium.";
RL Genes Dev. 8:2086-2096(1994).
CC -! FUNCTION: RECEPTOR FOR CAMP. REGULATES AXIAL PATTERNING AND
CC CELLULAR DIFFERENTIATION DURING LATE DEVELOPMENT. THE ACTIVITY OF
CC THIS RECEPTOR IS MEDIATED BY G PROTEINS.
CC -! SUBCELLULAR LOCATION: Integral membrane protein.
CC -! DEVELOPMENTAL STAGE: INITIALLY EXPRESSED DURING TIP ELONGATION AND
CC CONTINUES TO ACCUMULATE INTO CULMINATION.
CC -! PTM: CARBOXYL-TERMINAL SER OR THR RESIDUES MAY BE PHOSPHORYLATED.
CC -! SIMILARITY: BELONGS TO FAMILY 5 OF G-PROTEIN COUPLED RECEPTORS.
CC GCRDB; GCR_0277; -.
DR DictyDb; D00277; card.
DR InterPro; IPR000848; GPCR_CAMP.
DR InterPro; IPR000832; GPCR_secretin.
DR Pfam; PF00002; 7tm_2; 1.
DR PRINTS; PR00247; GPCR_CAMP.
DR PROSITE; PS50261; G_PROTEIN_RECP_F2_4; 1.
KW G-protein coupled receptor; Transmembrane; Glycoprotein;
KW Phosphorylation; Multigene family.
FT DOMAIN 1 11 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 12 32 1 (POTENTIAL).
FT DOMAIN 33 44 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 45 65 2 (POTENTIAL).
FT DOMAIN 66 89 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 90 110 3 (POTENTIAL).
FT DOMAIN 111 119 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 120 140 4 (POTENTIAL).
FT DOMAIN 141 161 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 162 182 5 (POTENTIAL).
FT DOMAIN 183 208 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 209 229 6 (POTENTIAL).
FT DOMAIN 230 263 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 264 284 7 (POTENTIAL).
FT DOMAIN 285 443 CYTOPLASMIC (POTENTIAL).
FT DOMAIN 293 443 ASN-RICH.
FT DOMAIN 293 298 POLY-ASN.
FT DOMAIN 333 340 POLY-GLN.
FT DOMAIN 343 353 POLY-GLN.
FT DOMAIN 412 426 POLY-ASN.
SQ SEQUENCE 443 AA; 51456 MW; CDF3A9DEE5A5BBE2 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 443;
Best Local Similarity 66.7%; Pred. No. 3.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
DB 72 EAGST 77

RESULT 11
VGLX_HSVBS STANDARD; PRT; 444 AA.
ID VGLX_HSVBS
AC Q08103;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Glycoprotein GX precursor.
OS Bovine herpesvirus type 1.2 (strain ST).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
OX NCBI_TaxID=45407;
RN [1]
SEQUENCE FROM N.A.
RX MEDLINE=94167875; PubMed=8122370;
RA Leung-Tack P., Audonnet J.F., Riviere M.;
RT "The complete DNA sequence and the genetic organization of the short
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RT unique region (US) of the bovine herpesvirus type 1 (ST strain).";
RL Virology 199:409-421(1994).
CC -----
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CC -----
CC EMBL; Z23068; CAA80603.1; -.
CC InterPro; IPR003363; Herpes_gg.
CC Pfam; PF02400; Herpes_gg; 1.
CC Glycoprotein; Transmembrane; Signal.
CC SIGNAL 1 24 POTENTIAL.
CC FT CHAIN 25 444 GLYCOPROTEIN GX.
CC FT TRANSMEM 390 414 POTENTIAL.
CC FT CARBOHYD 117 117 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 240 240 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 335 335 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC SEQUENCE 444 AA; 46708 MW; 0145942AA35B05CB CRC64;
CC -----
Query Match 90.5%; Score 19; DB 1; Length 444;
Best Local Similarity 66.7%; Pred. No. 3.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 eagxxs 6
Db 267 EAGSAS 272
ID YU88_MYCTU STANDARD; PRT; 474 AA.
AC OS3305;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hypothetical 50.9 kDa protein RV3088.
GN RV3088 OR MT3173 OR MV013.09.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae; Mycobacterium.
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RX MEDLINE=96295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekai F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., DeBoy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
RA Kolony J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W.;
RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO THE UPF0089 FAMILY.
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CC -----
CC EMBL; AL021309; CAA16146.1; -.
CC EMBL; AE007134; AAK47509.1; -.
CC TIGR; MT3173; -.
CC TubercuList; RV3088; -.
CC InterPro; IPR004255; UPF0089.
CC Pfam; PF03007; UPF0089; 1.
CC KW Hypothetical protein; Complete proteome.
CC SEQUENCE 474 AA; 50886 MW; 36832D972BE3851A CRC64;
CC -----
Query Match 90.5%; Score 19; DB 1; Length 474;
Best Local Similarity 66.7%; Pred. No. 3.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 eagxxs 6
Db 460 EAGTTS 465
ID NODT_RHILV STANDARD; PRT; 482 AA.
AC P15727;
DT 01-APR-1990 (Rel. 14, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Nodulation protein T precursor.
GN NODT.
OS Rhizobium leguminosarum (biovar viciae).
OC Plasmid sym PRLJ1.
OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
OC Rhizobiaceae; Rhizobium.
OX NCBI_TaxID=387;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=8401;
RX MEDLINE=90151607; PubMed=2303029;
RA Economou A., Hamilton W.D.O., Johnston A.W.B., Downie J.A.;
RA "The Rhizobium nodulation gene nodO encodes a Ca2(+)-binding protein
RT that is exported without N-terminal cleavage and is homologous to
RL haemolysin and related proteins.";
RL EMBO J. 9:349-354(1990).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=90251164; PubMed=2338917;
RA Surin B.P., Watson J.M., Hamilton W.D.O., Economou A., Downie J.A.;
RA "Molecular characterization of the nodulation gene, nodT, from two
RT biovars of Rhizobium leguminosarum.";
RL Mol. Microbiol. 4:245-252(1990).
CC -1- SUBCELLULAR LOCATION: Attached to the membrane by a lipid anchor
CC (Potential).
CC -1- SIMILARITY: BELONGS TO THE FUSA/NODT FAMILY.
CC -----
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CC -----
CC EMBL; X17285; CAA35177.1; -.
CC PIR; S08384; S08384.
CC PIR; S10133; S10133.
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DR InterPro; IPR000815; Hg_reductase.
DR InterPro; IPR001100; pyr_redox.
DR InterPro; IPR004099; pyr_redox_dim.
DR Pfam; PF00403; HMA; 1.
DR Pfam; PF00070; pyr_redox; 1.
DR Pfam; PF02852; pyr_redox_dim; 1.
DR PRINTS; PR00368; FADPNR.
DR PRINTS; PR00945; HGRDRTASEI.
DR PRINTS; PR00411; PNDRDTASEI.
DR PROSITE; PS01047; HMA_1; 1.
DR PROSITE; PS00846; HMA_2; 1.
DR PROSITE; PS00076; PYRIDINE_REDOX_1; 1.
KW Mercuric resistance; Oxidoreductase; Flavoprotein; FAD; NADP;
KW Mercuric; Redox-active center; Metal-binding; Plasmid.
FT DOMAIN 1 66 HMA.
FT NP_BIND 100 130 FAD (ADP PART) (PROBABLE).
FT DISULFID 136 141 REDOX-ACTIVE.
FT NP_BIND 393 403 FAD (FLAVIN PART) (BY SIMILARITY).
FT METAL 558 559 HG(2+) (POTENTIAL).
FT METAL 559 559 HG(2+) (POTENTIAL).
SQ SEQUENCE 561 AA; 58558 MW; 111E02A702C157D6 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 561;
Best Local Similarity 66.7%; Pred. No. 4.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
Db 45 EAGTSS 50

RESULT 16
MERA_ENTAG STANDARD; PRT; 561 AA.
AC P94702;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Mercuric reductase (EC 1.16.1.1) (Hg(II) reductase).
GN MERA.
OS Enterobacter agglomerans (Pantoea agglomerans).
OC Plasmid pKLH272.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Pantoea.
OX NCBI_TaxID=549;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97303086; PubMed=9159519;
RA Yurieva O., Kholodil G., Minakhin L., Gorlenko Z., Kalyaeva E.,
RA Mindlin S., Nikiforov V.;
RT "Intercontinental spread of promiscuous mercury-resistance
transposons in environmental bacteria.";
RL Mol. Microbiol. 24:321-329(1997).
CC -1- FUNCTION: RESISTANCE TO HG(2+) IN BACTERIA APPEARS TO BE GOVERNED
CC BY A SPECIALIZED SYSTEM WHICH INCLUDES MERCURIC REDUCTASE. MERA
CC PROTEIN IS RESPONSIBLE FOR VOLATILIZING MERCURY AS HG(0).
CC -1- CATALYTIC ACTIVITY: Hg + NADP(+) + H(+) = Hg(2+) + NADPH.
CC -1- COFACTOR: FAD.
CC -1- SUBUNIT: HOMODIMER (BY SIMILARITY).
CC -1- MISCELLANEOUS: THE ACTIVE SITE IS A REDOX-ACTIVE DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE PYRIDINE NUCLEOTIDE-DISULFIDE
CC OXIDOREDUCTASES CLASS-I.
CC -1- SIMILARITY: CONTAINS 1 HMA DOMAIN.
CC -----
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CC -----

EMBL; Y08992; CAA70184.1; -.
DR HSSP; Q04656; 1AW0.
DR InterPro; IPR001327; FAD_pyr_redox.
DR InterPro; IPR001934; HMA.
DR InterPro; IPR000815; Hg_reductase.
DR InterPro; IPR000205; NAD_binding.
DR InterPro; IPR001100; pyr_redox.
DR InterPro; IPR004099; pyr_redox_dim.
DR Pfam; PF00403; HMA; 1.
DR Pfam; PF00070; pyr_redox; 1.
DR Pfam; PF02852; pyr_redox_dim; 1.
DR PRINTS; PR00368; FADPNR.
DR PRINTS; PR00945; HGRDRTASEI.
DR PRINTS; PR00411; PNDRDTASEI.
DR PROSITE; PS01047; HMA_1; 1.
DR PROSITE; PS00846; HMA_2; 1.
DR PROSITE; PS00076; PYRIDINE_REDOX_1; 1.
KW Mercuric resistance; Oxidoreductase; Flavoprotein; FAD; NADP;
KW Mercuric; Redox-active center; Metal-binding; Plasmid.
FT DOMAIN 1 66 HMA.
FT NP_BIND 100 130 FAD (ADP PART) (PROBABLE).
FT DISULFID 136 141 REDOX-ACTIVE.
FT NP_BIND 393 403 FAD (FLAVIN PART) (BY SIMILARITY).
FT METAL 558 559 HG(2+) (POTENTIAL).
FT METAL 559 559 HG(2+) (POTENTIAL).
SQ SEQUENCE 561 AA; 58785 MW; FABA07D7EC2F13C8 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 561;
Best Local Similarity 66.7%; Pred. No. 4.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
Db 45 EAGTSS 50

RESULT 17
AC22_STRCO STANDARD; PRT; 578 AA.
ID AC22_STRCO
AC P46105;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Probable actinorhodin transporter.
GN ACT11-2 OR SCBAC28G1.09.
OS Streptomyces coelicolor.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces.
OX NCBI_TaxID=1902;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91347376; PubMed=1878971;
RA Fernandez-Moreno M.A., Caballero J.L., Hopwood D.A., Malpartida F.;
RT "The act cluster contains regulatory and antibiotic export genes,
direct targets for translational control by the bida trna gene of
Streptomyces.";
RN Cell 66:769-780(1991).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-A3(2);
RA Warren T., Harris D., Cerdeno A.M., Parkhill J., Barrell B.G.,
RA Rajandream M.A.;
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: PROMOTES THE EFFLUX OF ACTINORHODIN.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- SIMILARITY: BELONGS TO THE MAJOR FACILITATOR FAMILY (ALSO KNOWN
CC AS THE DRUG RESISTANCE TRANSLUCASE FAMILY).
CC -----
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DR EMBL: M64683; AAA26690.1; -
 DR EMBL: AL593842; CAC44196.1; -
 DR InterPro: IPR003662; subtransporter.
 DR Pfam: PF00083; sugar_tr; 1.
 KW Antibiotic resistance; Transport; Transmembrane.
 FT TRANSMEM 78 98 POTENTIAL.
 FT TRANSMEM 109 129 POTENTIAL.
 FT TRANSMEM 135 155 POTENTIAL.
 FT TRANSMEM 170 190 POTENTIAL.
 FT TRANSMEM 202 222 POTENTIAL.
 FT TRANSMEM 232 252 POTENTIAL.
 FT TRANSMEM 259 279 POTENTIAL.
 FT TRANSMEM 306 326 POTENTIAL.
 FT TRANSMEM 341 361 POTENTIAL.
 FT TRANSMEM 369 389 POTENTIAL.
 FT TRANSMEM 444 464 POTENTIAL.
 FT TRANSMEM 546 566 POTENTIAL.
 SQ SEQUENCE 578 AA; 59772 MW; E6C1DC75E6038B92 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 578;
 Best Local Similarity 66.7%; Pred. No. 4.8e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxvs 6
 |||||
 Db 430 EAGTAS 435

RESULT 18
 DNK2_SYNY3 STANDARD; PRT; 636 AA.
 AC P22358;
 DT 01-AUG-1991 (Rel. 19, Created)
 DT 01-AUG-1991 (Rel. 19, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Chapterone protein dnaK2 (Heat shock protein 70-2) (Heat shock 70 kDa
 DE protein 2) (HSP70-2).
 DE DnaK2 OR DnaK OR SLL0170.
 GN Synchocystis sp. (strain PCC 6803).
 OS Bacteria; Cyanobacteria; Chroococcales; Synchocystis.
 OC NCBI_TaxID=1148;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91093185; PubMed=1670771;
 RA Chitnis P.R., Nelson N.;
 RT "Molecular cloning of the genes encoding two chaperone proteins of
 RT the cyanobacterium Synchocystis sp. PCC 6803.";
 RL J. Biol. Chem. 266:58-65(1991).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=96127529; PubMed=8590279;
 RA Kaneko T., Tanaka A., Sato S., Kotani H., Sazuka T., Miyajima N.,
 RA Sugitara M., Tabata S.;
 RT "Sequence analysis of the genome of the unicellular cyanobacterium
 RT Synchocystis sp. strain PCC6803. I. Sequence features in the 1 Mb
 RT region from map positions 64% to 92% of the genome.";
 RL DNA Res. 2:153-166(1995).
 CC -!- FUNCTION: ACTS AS A CHAPERONE (BY SIMILARITY).
 CC -!- INDUCTION: BY STRESS CONDITIONS E.G. HEAT SHOCK.
 CC -!- SIMILARITY: BELONGS TO THE HEAT SHOCK PROTEIN 70 FAMILY.
 CC
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CC -----
 DR EMBL: M57518; AAA27287.1; -
 DR EMBL: D63999; BAA10059.1; -
 DR PIR: C39025; G39025.
 DR HSSP: P04475; 1DG4.
 DR InterPro: IPR001023; HSP70.
 DR Pfam: PF00012; HSP70; 1.
 DR PRINTS: PR00301; HEATSHOCK70.
 DR PROSITE: PS00297; HSP70.1; 1.
 DR PROSITE: PS00329; HSP70_2; 1.
 DR PROSITE: PS01036; HSP70_3; 1.
 KW Chaperone; ATP-binding; Heat shock; Multigene family;
 KW Complete proteome.
 SQ SEQUENCE 636 AA; 67614 MW; 33AE4CECBA28F40A CRC64;

Query Match 90.5%; Score 19; DB 1; Length 636;
 Best Local Similarity 66.7%; Pred. No. 5.2e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxvs 6
 |||||
 Db 615 EAGTSS 620

RESULT 19
 TOXA_PSEAE STANDARD; PRT; 638 AA.
 AC P11439; Q91417;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Exotoxin A precursor (NAD-dependent ADP-ribosyltransferase
 DE (EC 2.4.2.-)).
 GN ETA OR P1148.
 OS Pseudomonas aeruginosa.
 OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;
 OC Pseudomonas.
 OX NCBI_TaxID=287;
 RN [1]
 RP SEQUENCE FROM N.A., AND SEQUENCE OF 26-53.
 RX MEDLINE=84194063; PubMed=6201861;
 RA Gray G.L., Heyneker H.L.;
 RA Chen E.Y., Smith D.H., Balbridge J.S., Harkins R.N., Vasil M.L.,
 RT "Cloning, nucleotide sequence, and expression in Escherichia coli of
 RT the exotoxin A structural gene of Pseudomonas aeruginosa.";
 RL Proc. Natl. Acad. Sci. U.S.A. 81:2645-2649(1984).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX STRAIN=ATCC 15692 / PA01;
 RC MEDLINE=20437337; PubMed=10984043;
 RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
 RA Hickey M.J., Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,
 RA Garber R.L., Goltry L., Tolentino E., Westbrock-Wadman S., Yuan Y.,
 RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
 RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
 RA Reizer J., Sailer M.H., Hancock R.E.W., Lory S., Olson M.V.;
 RT "Complete genome sequence of Pseudomonas aeruginosa PA01, an
 RT opportunistic pathogen.";
 RL Nature 406:959-964(2000).
 RN [3]
 RP ACTIVE SITE.
 RX MEDLINE=87250491; PubMed=2885323;
 RA Carroll S.F., Collier R.J.;
 RT "Active site of Pseudomonas aeruginosa exotoxin A. Glutamic acid 553
 RT is photolabeled by NAD and shows functional homology with glutamic
 RT acid 148 of diphtheria toxin.";
 RL J. Biol. Chem. 262:8707-8711(1987).
 RN [4]
 RP DOMAINS.
 RX MEDLINE=90375493; PubMed=2118903;
 RA Chaudhary V.K., Jinno Y., Galo M.G., Fitzgerald D., Pastan I.;
 RT "Mutagenesis of Pseudomonas exotoxin in identification of sequences

responsible for the animal toxicity.";
J. Biol. Chem. 265:16306-16310(1990).
[5]
RP DOMAINS.
RX MEDLINE=91006124; PubMed=2170123;
RA Bourdenet S., Vacheron M.-J., Guinand M., Michel G., Arminjon F.;
RT "Biochemical and immunochemical studies of proteolytic fragments of
RT exotoxin A from Pseudomonas aeruginosa.";
RL Eur. J. Biochem. 192:379-385(1990).
[6]
RX X-RAY CRYSTALLOGRAPHY (3.0 ANGSTROMS) OF 424-638.
RA MEDLINE=96016159; PubMed=7568123;
RX Li M., Dyda F., Benhar I., Pastan I., Davies D.R.;
RT "The crystal structure of Pseudomonas aeruginosa exotoxin domain III
RT with nicotinamide and AMP: conformational differences with the intact
RT exotoxin.";
RL Proc. Natl. Acad. Sci. U.S.A. 92:9308-9312(1995).
[7]
RX X-RAY CRYSTALLOGRAPHY (2.3 ANGSTROMS) OF 424-638.
RA MEDLINE=96293446; PubMed=8692916;
RX Li M., Dyda F., Benhar I., Pastan I., Davies D.R.;
RT "Crystal structure of the catalytic domain of Pseudomonas exotoxin A
RT complexed with a nicotinamide adenine dinucleotide analog:
RT implications for the activation process and for ADP ribosylation.";
RL Proc. Natl. Acad. Sci. U.S.A. 93:6902-6906(1996).
CC -1- FUNCTION: THIS TOXIN IS AN NAD-DEPENDENT ADP-RIBOSYLTRANSFERASE.
CC IT CATALYZES THE TRANSFER OF THE ADP RIBOSYL MOIETY OF OXIDIZED
CC NAD ONTO ELONGATION FACTOR 2 (EF-2) THUS ARRESTING PROTEIN
CC SYNTHESIS.
CC -1- PTM: THE 8 CYSTEINES PARTICIPATE IN INTRACHAIN DISULFIDE BONDS.
CC -1- SIMILARITY: REGIONAL SEQUENCE SIMILARITY AT THE ACTIVE SITE
CC WITH DIPHTHERIA TOXIN (DT).
CC
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CC
CC EMBL; K01397; AAB59097.1; -;
CC PIR; A30347; A30347.
CC PDB; 1AER; 10-JUN-96.
CC PDB; 1DMA; 15-SEP-95.
CC Toxin; Signal; Transferase; Glycosyltransferase; NAD; 3D-structure;
CC Complete proteome.
FT SIGNAL 1 25
FT CHAIN 26 638 EXOTOXIN A.
FT DOMAIN 26 277 IA (REQUIRED FOR TARGET CELL
FT RECOGNITION).
FT DOMAIN 278 389 II (REQUIRED FOR TRANSLLOCATION IN TARGET
FT CELL CYTOPLASM).
FT DOMAIN 390 429 IB.
FT DOMAIN 430 638 III (REQUIRED FOR ADP-RIBOSYL ACTIVITY).
FT ACT_SITE 465 465 INTERACT WITH NAD.
FT CONFLICT 4 4 T -> I (IN REF. 1).
FT CONFLICT 22 22 F -> S (IN REF. 1).
FT CONFLICT 204 204 A -> T (IN REF. 1).
FT CONFLICT 389 389 S -> N (IN REF. 1).
FT CONFLICT 432 432 I -> V (IN REF. 1).
FT CONFLICT 540 540 G -> S (IN REF. 1).
SQ SEQUENCE 638 AA; 69284 MW; 7B9AAD56A27C700A CRC64;

Query Match 90.5%; Score 19; DB 1; Length 638;
Best Local Similarity 66.7%; Pred. No. 5.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |

Db 384 EAGAAS 389
RESULT 20
SGT1_ARATH
ID SGT1_ARATH STANDARD; PRT; 643 AA.
AC Q9LSM5;
DT 01-MAR-2002 (Rel. 41, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE SGT1 protein homolog At5g65490.
GN AF5G65490 OR K1904.2.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. COLUMBIA;
RA Kaneko T., Katoh T., Asamizu E., Sato S., Nakamura Y., Kotani H.,
RA Tabata S.;
RT "Structural analysis of Arabidopsis thaliana chromosome 5. XI.";
RL Submitted (APR-1999) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO THE SGT1 FAMILY.
CC
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CC
CC EMBL; AB026638; BAA98169.1; -;
KW Hypothetical protein.
SQ SEQUENCE 643 AA; 73161 MW; 9F9DD990F65B4C4F CRC64;

Query Match 90.5%; Score 19; DB 1; Length 643;
Best Local Similarity 66.7%; Pred. No. 5.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |

Db 445 EAGSSS 450

RESULT 21
EOMD_XENLA
ID EOMD_XENLA STANDARD; PRT; 692 AA.
AC P79944;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Eomesodermin.
GN EOMES.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidea; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Gastrula;
RA MEDLINE=97133207; PubMed=8978604;
RT Ryan K., Garrett N., Mitchell A., Gurdon J.B.;
RL "Eomesodermin, a key early gene in Xenopus mesoderm differentiation.";
RL Cell 87:989-1000(1996).
CC -1- FUNCTION: INVOLVED IN MESODERM DIFFERENTIATION. ACTIVATES WNT8,
CC BRACHYURY, CHD AND MIX.1 EXPRESSION.
CC -1- SUBCELLULAR LOCATION: Nuclear (Potential).
CC -1- DEVELOPMENTAL STAGE: FIRST EXPRESSED AT OR JUST AFTER MIDBLASTULA

CC TRANSITION (STAGE 8). MAXIMALLY EXPRESSED AT STAGE 10 AS AN
CC EQUITATORIAL MESODERM BAND, MORE PROMINENTLY ON THE DORSAL SIDE
CC AND AROUND THE INVAGINATING DORSAL LIP.
CC -!- INDUCTION: BY ACTIVIN.
CC -!- DOMAIN: CONTAINS 13 S-P-X-X REPEATS.
CC -!- SIMILARITY: CONTAINS 1 T-BOX DOMAIN.
CC
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CC
CC EMBL; U75996; AAC60061.1; -
CC HSSP; P24781; 1XBR.
CC InterPro; IPR001699; T-box.
CC Pfam; PF00907; T-box; 1.
CC PRINTS; PR00937; TBOX.
CC SMART; SM00425; TBOX; 1.
CC PROSITE; PS01283; TBOX_1; 1.
CC PROSITE; PS01264; TBOX_2; 1.
CC PROSITE; PS02522; TBOX_3; 1.
CC Developmental protein; Transcription regulation; DNA-binding;
KW Nuclear protein; Repeat.
FT DNA_BIND 263 443
FT SEQUENCE 692 AA; 75943 MW; 9D129A67F6357989 CRC64;
SQ
Query Match 90.5%; Score 19; DB 1; Length 692;
Best Local Similarity 66.7%; Pred. No. 5.7e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 eaqxss 6
DB 90 EAGSSS 95
RESULT 22
PCGB_MOUSE STANDARD; PRT; 883 AA.
AC Q61361;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 30-NOV-2000 (Rel. 39, Last annotation update)
DE Brevican core protein precursor.
GN BCAN.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-BALB/C; TISSUE=Brain;
RX MEDLINE=97432816; PubMed=9286696;
RA Rauch U., Meyer H., Brakebusch C., Seidenbecher C., Gundelfinger E.D.,
RA Beier D.R., Fassler R.;
RT "Sequence and chromosomal localization of the mouse brevican gene.";
RL Genomics 44:15-21(1997).
CC -!- FUNCTION: MAY PLAY A ROLE IN THE TERMINALLY DIFFERENTIATING AND
CC THE ADULT NERVOUS SYSTEM DURING POSTNATAL DEVELOPMENT. COULD
CC STABILIZE INTERACTIONS BETWEEN HA AND BRAIN PROTEOGLYCAN.
CC -!- SUBCELLULAR LOCATION: SECRETED; EXTRACELLULAR MATRIX (BY
CC SIMILARITY).
CC -!- TISSUE SPECIFICITY: BRAIN (BY SIMILARITY).
CC -!- PTM: CONTAINS MOSTLY CHONDROITIN SULFATE (BY SIMILARITY).
CC -!- SIMILARITY: CONTAINS 1 IMMUNOGLOBULIN-LIKE V-TYPE DOMAIN.
CC -!- SIMILARITY: CONTAINS 2 LINK DOMAINS.
CC -!- SIMILARITY: CONTAINS 1 EGF-LIKE DOMAIN.
CC -!- SIMILARITY: CONTAINS 1 C-TYPE LECTIN FAMILY DOMAIN.
CC -!- SIMILARITY: CONTAINS 1 SUSHI (SCR) DOMAIN.
CC -!- SIMILARITY: BELONGS TO THE AGGREGAN/VERSICAN PROTEOGLYCAN FAMILY.

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CC
CC EMBL; X87096; CAA60575.1; -
CC HSSP; P20693; 1HLJ.
CC MGD; MGI:1096385; Bcan.
CC InterPro; IPR000561; EGF-like.
CC InterPro; IPR000742; EGF_2.
CC InterPro; IPR003006; Ig_MHC.
CC InterPro; IPR003596; Ig_v.
CC InterPro; IPR000538; Link.
CC InterPro; IPR000436; Sushi_SCR_CCP.
CC InterPro; IPR001304; Lectin_c.
CC Pfam; PF00008; EGF; 1.
CC Pfam; PF00047; ig; 1.
CC Pfam; PF00059; lectin_c; 1.
CC Pfam; PF00084; sushi; 1.
CC Pfam; PF00193; xlink; 2.
CC ProDom; PD000918; Link; 2.
CC SMART; SM00032; CCP; 1.
CC SMART; SM00034; CLECT; 1.
CC SMART; SM00181; EGF; 1.
CC SMART; SM00406; IGV; 1.
CC SMART; SM00445; LINK; 2.
CC PROSITE; PS00022; EGF_1; 1.
CC PROSITE; PS01186; EGF_2; 1.
CC PROSITE; PS00290; IG_MHC; 1.
CC PROSITE; PS01241; LINK; 2.
CC PROSITE; PS00615; C-TYPE-LECTIN_1; 1.
CC PROSITE; PS00641; C-TYPE-LECTIN_2; 1.
CC Glycoprotein; Hyaluronic acid; Proteoglycan; Lectin; Signal; Sushi;
KW EGF-like domain; Repeat; Immunoglobulin domain.
FT SIGNAL 1 22
FT CHAIN 23 883
FT DOMAIN 32 157
FT DOMAIN 173 250
FT DOMAIN 271 352
FT DOMAIN 622 658
FT DOMAIN 658 786
FT DOMAIN 787 851
FT DISULFID 56 136
FT DISULFID 178 249
FT DISULFID 202 223
FT DISULFID 276 351
FT DISULFID 300 321
FT DISULFID 626 637
FT DISULFID 631 646
FT DISULFID 648 657
FT DISULFID 664 675
FT DISULFID 692 784
FT DISULFID 760 776
FT DISULFID 791 834
FT DISULFID 820 847
FT CARBOHYD 129 129
FT CARBOHYD 336 336
SQ SEQUENCE 883 AA; 96013 MW; CC2C3C97B453B45 CRC64;
Query Match 90.5%; Score 19; DB 1; Length 883;
Best Local Similarity 66.7%; Pred. No. 7.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 eaqxss 6
DB 558 EAGSSS 563

RESULT 23

PCCB_RAT STANDARD: PRT; 883 AA.
AC P55068; Q63040; Q62860; Q63513;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Brevican core protein precursor (Brain enriched hyaluronan binding protein) (BEHAB protein).
GN BCAN.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RN SEQUENCE FROM N.A.
RC STRAIN=SPRAGUE-DAWLEY; TISSUE=Brain;
RX MEDLINE=96070828; PubMed=7592978;
RA Seidenbecher C.L., Richter K., Rauch U., Faessler R., Garner C.C., Gundelfinger E.D.;
RT "Brevican, a chondroitin sulfate proteoglycan of rat brain, occurs as secreted and cell surface glycosylphosphatidylinositol-anchored isoforms.";
RL J. Biol. Chem. 270:27206-27212(1995).
RN [2]
RN SEQUENCE FROM N.A., AND SEQUENCE OF 396-407.
RC TISSUE=Brain;
RX MEDLINE=96074575; PubMed=7488217;
RA Yamada H., Watanabe K., Shimonaka M., Yamasaki M., Yamaguchi Y.;
RT "cDNA cloning and the identification of an aggrecanase-like cleavage site in rat brevican.";
RL Biochem. Biophys. Res. Commun. 216:957-963(1995).
RN [3]
RN SEQUENCE OF 1-423 FROM N.A.
RC STRAIN=SPRAGUE-DAWLEY; TISSUE=Brain;
RX MEDLINE=94216386; PubMed=7512973;
RA Jaworski D.M., Kelly G.M., Hockfield S.;
RT "BEHAB, a new member of the proteoglycan tandem repeat family of hyaluronan-binding proteins that is restricted to the brain.";
RL J. Cell Biol. 125:495-509(1994).
CC -!- FUNCTION: MAY PLAY A ROLE IN THE TERMINALLY DIFFERENTIATING AND THE ADULT NERVOUS SYSTEM DURING POSTNATAL DEVELOPMENT. COULD STABILIZE INTERACTIONS BETWEEN HA AND BRAIN PROTEOGLYCANS. THE GPI-ANCHORED ISOFORM MAY FUNCTION AS A CHONDROITIN SULFATE-BEARING CELL SURFACE RECEPTOR.
CC -!- SUBCELLULAR LOCATION: SECRETED; EXTRACELLULAR MATRIX AND ONE FORM ATTACHED TO THE MEMBRANE BY A GPI-ANCHOR.
CC -!- ALTERNATIVE PRODUCTS: 2 ISOFORMS: A SECRETED FORM (SHOWN HERE) AND A GPI-ANCHORED FORM; ARE PRODUCED BY ALTERNATIVE SPLICING.
CC -!- TISSUE SPECIFICITY: BRAIN.
CC -!- DEVELOPMENTAL STAGE: SOLUBLE FORM INCREASES FROM DAY P4 TO P64. GPI-ANCHORED ISOFORM INCREASES AFTER DAY P8.
CC -!- PTM: CONTAINS MOSTLY CHONDROITIN SULFATE.
CC -!- SIMILARITY: CONTAINS 1 IMMUNOGLOBULIN-LIKE V-TYPE DOMAIN.
CC -!- SIMILARITY: CONTAINS 2 LINK DOMAINS.
CC -!- SIMILARITY: CONTAINS 1 EGF-LIKE DOMAIN.
CC -!- SIMILARITY: CONTAINS 1 C-TYPE LECTIN FAMILY DOMAIN.
CC -!- SIMILARITY: CONTAINS 1 SUSHI (SCR) DOMAIN.
CC -!- SIMILARITY: BELONGS TO THE AGGREGAN/VERSICAN PROTEOGLYCAN FAMILY.
CC -!- CAUTION: REF.3 SEQUENCE DIFFERS FROM THAT SHOWN FROM POSITION 364 ONWARD AND IS SMALLER (371 AA) DUE TO A FRAMESHIFT.
CC -----
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CC -----
CC EMBL; X79881; CAA56255.1; -;
CC EMBL; X85406; CAA60160.1; -;
CC EMBL; U37142; AAA87847.1; -;

DR EMBL; Z28366; CAA82215.1; ALT_FRAME.
DR HSP; P20693; IHLJ.
DR InterPro; IPR000561; EGF-like.
DR InterPro; IPR000742; EGF_2.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR003596; Ig_v.
DR InterPro; IPR000538; Link.
DR InterPro; IPR000436; Sushi_SCR_CCP.
DR InterPro; IPR001304; lectin_c.
DR Pfam; PF00008; EGF; 1.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00059; lectin_c; 1.
DR Pfam; PF00084; sushi; 1.
DR Pfam; PF00193; Xlink; 2.
DR ProDom; PD000918; Link; 2.
DR SMART; SM00032; CCP; 1.
DR SMART; SM00034; CLECT; 1.
DR SMART; SM00181; EGF; 1.
DR SMART; SM00406; IGV; 1.
DR SMART; SM00445; Link; 2.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR PROSITE; PS01241; LINK; 2.
DR PROSITE; PS00615; C-TYPE_LECTIN_1; 1.
DR PROSITE; PS00041; C-TYPE_LECTIN_2; 1.
DR KW Glycoprotein; Hyaluronic acid; Proteoglycan; Lectin; Signal; Sushi;
DR KW EGF-like domain; Repeat; Immunoglobulin domain; Alternative splicing;
DR GPI-anchor.
FT SIGNAL 1 22 POTENTIAL.
FT CHAIN 23 883 BREVICAN CORE PROTEIN.
FT DOMAIN 32 157 IG-LIKE V-TYPE DOMAIN.
FT DOMAIN 173 250 LINK 1.
FT DOMAIN 271 352 LINK 2.
FT DOMAIN 622 658 EGF-LIKE.
FT DOMAIN 658 786 C-TYPE LECTIN.
FT DOMAIN 787 851 SUSHI.
FT DISULFID 56 136 BY SIMILARITY.
FT DISULFID 178 249 BY SIMILARITY.
FT DISULFID 202 223 BY SIMILARITY.
FT DISULFID 276 351 BY SIMILARITY.
FT DISULFID 300 321 BY SIMILARITY.
FT DISULFID 626 637 BY SIMILARITY.
FT DISULFID 631 646 BY SIMILARITY.
FT DISULFID 648 657 BY SIMILARITY.
FT DISULFID 791 834 BY SIMILARITY.
FT DISULFID 820 847 BY SIMILARITY.
FT CARBOHYD 129 129 N-LINKED (GLCNAC...) (POTENTIAL).
FT CARBOHYD 336 336 N-LINKED (GLCNAC...) (POTENTIAL).
FT VARSPLIC 625 645 DCIFSPCHNGTCLKEGFR -> NSAEGSNPAFLFLLL
FT VARSPLIC 646 883 OLMDT (IN GPI-ANCHORED ISOFORM).
FT VARSPLIC 51 52 MISSING (IN GPI-ANCHORED ISOFORM).
FT CONFLICT 503 503 AL -> WV (IN REF. 3).
FT CONFLICT 518 519 TV -> PA (IN REF. 2).
FT CONFLICT 526 526 G -> R (IN REF. 2).
FT CONFLICT 541 541 G -> A (IN REF. 2).
FT CONFLICT 556 556 R -> S (IN REF. 2).
FT CONFLICT 573 573 E -> A (IN REF. 2).
FT CONFLICT 583 583 V -> L (IN REF. 2).
FT CONFLICT 649 649 V -> L (IN REF. 2).
FT CONFLICT 670 670 P -> A (IN REF. 2).
FT CONFLICT 738 738 P -> A (IN REF. 2).
FT CONFLICT 809 809 R -> A (IN REF. 2).
SQ SEQUENCE 883 AA; 96057 MW; AC7ACC40CB53ED37 CRC64;

Query Match 90.58; Score 19; DB 1; Length 883;
Best Local Similarity 66.7%; Pred. No. 7.3e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
||| |

Db 558 EAGSSS 563

RESULT 24
SRCA_RABIT
ID SRCA_RABIT STANDARD; PRT; 908 AA.
AC P13666;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Sarcalumenin precursor.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=89345602; PubMed=2762314;
RA Leberer E., Charuk J.H.M., Green N.M., MacLennan D.H.;
RT "Molecular cloning and expression of cDNA encoding a lumenal calcium
binding glycoprotein from sarcoplasmic reticulum.";
RL Proc. Natl. Acad. Sci. U.S.A. 86:6047-6051(1989).
RN [2]
RP SEQUENCE OF 1-19 AND 458-908 FROM N.A.
RX MEDLINE=89123480; PubMed=2521635;
RA Leberer E., Charuk J.H.M., Clarke D.M., Green N.M., Zubrycka-Gaarn E.,
RA MacLennan D.H.;
RT "Molecular cloning and expression of cDNA encoding the 53,000-dalton
glycoprotein of rabbit skeletal muscle sarcoplasmic reticulum.";
RL J. Biol. Chem. 264:3484-3493(1989).
CC -|- FUNCTION: PERHAPS INVOLVED IN THE REGULATION OF CALCIUM TRANSPORT.
CC -|- SUBCELLULAR LOCATION: SARCOPLASMIC RETICULUM LUMEN. ASSOCIATED
CC THROUGH CA(2+) WITH THE MEMBRANE.
CC -|- ALTERNATIVE PRODUCTS: 2 ISOFORMS: A CALCIUM-BINDING
CC GLYCOPROTEIN/160 KDA (SHOWN HERE) AND A SECOND GLYCOPROTEIN/53
CC KDA; MAY BE PRODUCED BY ALTERNATIVE SPLICING.
CC
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CC
CC EMBL; M25750; AAA31189.1; -
DR EMBL; J04480; AA60730.1; -
DR PIR; A33280; A33280.
DR PIR; A33312; A33312.
KW Calcium-binding; Glycoprotein; Signal; Alternative splicing.
FT SIGNAL 1 19
FT CHAIN 20 908 160 KDA SARCALUMENIN.
FT CHAIN 458 908 53 KDA SARCALUMENIN.
FT DOMAIN 20 457 ACIDIC DOMAIN, PROBABLY BINDS CALCIUM.
FT CARBOHYD 102 102 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 716 716 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 824 824 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT VARSPLIC 21 456 MISSING (IN 53 KDA ISOFORM).
FT CONFLICT 474 474 O -> E (IN REF. 2).
FT CONFLICT 474 474 O -> E (IN REF. 2).
SQ SEQUENCE 908 AA; 97920 MW; A48CAA221AE1418B CRC64;

Query Match 90.5%; Score 19; DB 1; Length 908;
Best Local Similarity 66.7%; Pred. No. 7.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaaxxs 6
Db 397 EAGAAS 402
| | | | |

RESULT 25
POLS_IBDVA
ID POLS_IBDVA STANDARD; PRT; 1012 AA.
AC P08364;
DT 01-AUG-1988 (Rel. 08, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Structural polyprotein [Contains: Major structural protein VP2;
DE Nonstructural protein VP4; Minor structural protein VP3].
OS Avian infectious bursal disease virus (strain Australian 002-73)
OS (IBDV).
OC Viruses; dsRNA viruses; Birnaviridae; Avibirnavirus.
OX NCBI_TaxID=10997;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=86259073; PubMed=3014441;
RA Hudson P.J., McKern N.M., Power B.E., Azad A.A.;
RT "Genomic structure of the large RNA segment of infectious bursal
RT disease virus.";
RL Nucleic Acids Res. 14:5001-5012(1986).
RN [2]
RP SEQUENCE OF 703-1012 FROM N.A.
RX MEDLINE=86220784; PubMed=3011501;
RA Hudson P.J., McKern N.M., Fahey K.J., Azad A.A.;
RT "Predicted sequence of the host-protective immunogen of infectious
RT bursal disease virus.";
RL FEBS Lett. 201:143-146(1986).
CC -|- FUNCTION: SEGMENT A ENCODES A POLYPROTEIN, THAT IS PROCESSED INTO
CC THE MAJOR STRUCTURAL PROTEINS OF THE VIRION VP2 AND VP3, AND INTO
CC THE PUTATIVE PROTEASE VP4.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; X03993; CAA27629.1; ALT_INIT.
DR PIR; A24382; GNXSAU.
DR MEROPS; S50.002; -
DR InterPro; IPR002662; Birna_VP2.
DR InterPro; IPR002663; Birna_VP3.
DR InterPro; IPR002664; Birna_VP4.
DR Pfam; PF01766; Birna_VP2; 1.
DR Pfam; PF01767; Birna_VP3; 1.
DR Pfam; PF01768; Birna_VP4; 1.
KW Polyprotein; Structural protein; Nonstructural protein; Hydrolase;
KW Protease.
FT CHAIN 1 452 MAJOR STRUCTURAL PROTEIN VP2.
FT CHAIN 454 722 NONSTRUCTURAL PROTEIN VP4 (PROTEASE).
FT CHAIN 724 1012 MINOR STRUCTURAL PROTEIN VP3.
SQ SEQUENCE 1012 AA; 109503 MW; D9320A90459DE8F6 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1012;
Best Local Similarity 66.7%; Pred. No. 8.4e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaaxxs 6
Db 484 EAGAAS 489
| | | | |

RESULT 26
TSCC_HUMAN
ID TSCC_HUMAN STANDARD; PRT; 1021 AA.
AC P55017;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Thiazide-sensitive sodium-chloride cotransporter (NA-CL symporter).
GN SLC12A3 OR TSC.
OS Homo sapiens (Human).

CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A. AND VARIANTS GS.
 RX MEDLINE=96122035; PubMed=8528245;
 RA Simon D.B., Nelson-Williams C., Bia M.J., Ellison D., Karet F.E.,
 RA Molina A.M., Vawter I., Iwata F., Cushman H.M., Koolen M., Gainza F.J.,
 RA Gitelman H.J., Lifton R.P.;
 RT "Gitelman's variant of Bartter's syndrome, inherited hypokalaemic
 RT alkalosis, is caused by mutations in the thiazide-sensitive Na-Cl
 RT cotransporter.";
 RL Nat. Genet. 12:24-30(1996).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Kidney;
 RX MEDLINE=97001149; PubMed=8812482;
 RA Mastroianni N., de Fusco M., Zollo M., Arrigo G., Zuffardi O.,
 RA Bettinelli A., Ballabio A., Casari G.;
 RT "Molecular cloning, expression pattern, and chromosomal localization
 RT of the human Na-Cl thiazide-sensitive cotransporter (SLC12A3).";
 RL Genomics 35:486-493(1996).
 CC -!- FUNCTION: ELECTRICALLY SILENT TRANSPORTER SYSTEM WHICH IS A
 CC MEDIATOR OF SODIUM AND CHLORIDE REABSORPTION.
 CC -!- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -!- TISSUE SPECIFICITY: PREDOMINANT IN KIDNEY.
 CC -!- DISEASE: DEFECTS IN SLC12A3 ARE THE CAUSE OF GITELMAN'S SYNDROME
 CC (GS). AN AUTOSOMAL RECESSIVE DISEASE CHARACTERIZED BY DIVERSE
 CC ABNORMALITIES IN ELECTROLYTE HOMEOSTASIS INCLUDING HYPOKALAEMIC
 CC METABOLIC ALKALOSIS. GS IS A SUBSET OF BARTTER'S SYNDROME.
 CC -!- SIMILARITY: BELONGS TO THE SLC12A FAMILY OF TRANSPORTERS.
 CC -----
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 CC -----
 CC EMBL; U44128; AAC50355.1; -;
 DR EMBL; X91220; CAA62613.1; -;
 DR MM; 600968; -;
 DR MM; 263800; -;
 DR InterPro; IPR002293; AA_rel_permease.1.
 DR InterPro; IPR002948; NaCl transporter.
 DR PRINTS; PR01230; NACLTRANSPORT.
 KW Transport; Transmembrane; Glycoprotein; Disease mutation.
 FT DOMAIN 1 135 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 136 156 POTENTIAL.
 FT DOMAIN 157 158 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 159 179 POTENTIAL.
 FT DOMAIN 180 218 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 219 239 POTENTIAL.
 FT DOMAIN 240 261 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 262 282 POTENTIAL.
 FT DOMAIN 283 286 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 287 307 POTENTIAL.
 FT DOMAIN 308 339 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 340 360 POTENTIAL.
 FT DOMAIN 361 377 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 378 398 POTENTIAL.
 FT DOMAIN 399 452 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 453 473 POTENTIAL.
 FT DOMAIN 474 511 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 512 532 POTENTIAL.
 FT DOMAIN 533 534 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 535 555 POTENTIAL.
 FT DOMAIN 556 577 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 578 598 POTENTIAL.
 FT DOMAIN 599 660 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 661 681 POTENTIAL.
 FT DOMAIN 682 1021 CYTOPLASMIC (POTENTIAL).
 FT DOMAIN

FT CARBOHYD 406 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 426 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT VARIANT 209 R -> W (IN GS).
 FT VARIANT 349 /FTid=VAR_007113.
 FT VARIANT 421 /FTid=VAR_007114.
 FT VARIANT 486 /FTid=VAR_007115.
 FT VARIANT 496 /FTid=VAR_007116.
 FT VARIANT 561 /FTid=VAR_007117.
 FT VARIANT 588 /FTid=VAR_007118.
 FT VARIANT 630 /FTid=VAR_007119.
 FT VARIANT 655 /FTid=VAR_007120.
 FT VARIANT 655 /FTid=VAR_007121.
 FT VARIANT 728 /FTid=VAR_007122.
 FT VARIANT 741 /FTid=VAR_007123.
 FT VARIANT 850 /FTid=VAR_007124.
 FT VARIANT 955 /FTid=VAR_007125.
 FT CONFLICT 459 /FTid=VAR_007126.
 FT CONFLICT 766 AG -> VV (IN REF. 2).
 FT CONFLICT 807 E -> D (IN REF. 2).
 FT SEQUENCE 1021 AA; 113138 MW; D7ECE53DA6233821 CRC64;
 SQ
 QY 1 eagxxs 6
 Db 121 EAGTSS 126
 Query Match 90.5%; Score 19; DB 1; Length 1021;
 Best Local Similarity 66.7%; Pred. No. 8.5e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 RESULT 27
 ISWL_DROME
 ID ISWL_DROME STANDARD; PRT: 1027 AA.
 AC Q24368; Q9V6E8;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Iswi protein (imitation swi protein) (Nucleosome remodeling factor 140
 DE kda subunit) (NURF-140) (CHRC 140 kda subunit).
 GN ISWI OR CG8625.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ISO-1;
 RX MEDLINE=94187693; PubMed=7908117;
 RA Eifring L.K., Deuring R., McCallum C.M., Peterson C.L., Tamkun J.W.;
 RT "Identification and characterization of Drosophila relatives of the
 RL yeast transcriptional activator SNF2/SWI2.";
 RL Mol. Cell. Biol. 14:2225-2234(1994).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BERKELEY.1;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celnik S.E., Holt R.A., Evans C.A., Gocayne J.D.,

RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
RA Sutton G.G., Wortman J.R., Vandell M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
RA Wan K.H., Doyle C., Baxter E.G., Helt G.G., Nelson C.R., Miklos G.L.G.,
RA April J.F., Agbayani A., An H.-J., Andrews-Frankoch C., Baldwin D.,
RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brotter P.,
RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
RA de Pablo B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
RA Fostler C., Gabriellian A.E., Garg N.S., Gelbart W.M., Glasser K.,
RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
RA Jaitani M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RA "The genome sequence of *Drosophila melanogaster*.";
RL Science 287:2185-2195(2000).
CC -!- FUNCTION: COMPONENT OF THE NUCLEOSOME REMODELING FACTOR COMPLEX
CC (NURF), A PROTEIN COMPLEX THAT FACILITATES THE PERTURBATION OF
CC CHROMATIN STRUCTURE IN VITRO IN AN ATP-DEPENDENT MANNER. THE
CC HYDROLYSIS OF ATP DURING THE REMODELING OF CHROMATIN IS LIKELY TO
CC BE MEDIATED BY ISWI, RELEASING INORGANIC PHOSPHATE. IT IS ALSO A
CC COMPONENT OF THE ATP-UTILIZING CHROMATIN ASSEMBLY AND REMODELING
CC FACTOR (ACF) AND OF THE CHROMATIN ACCESSIBILITY COMPLEX (CHAC).
CC THIS SUBUNIT MAY SERVE AS THE ENERGY-TRANSDUCING COMPONENT OF
CC CHROMATIN-REMODELING MACHINES.
CC -!- SUBUNIT: NURF IS COMPOSED OF FOUR SUBUNITS: A 215 kDa PROTEIN,
CC IMITATION SWITCH (ISWI), NURF-55, AND NURF-38.
CC -!- SUBCELLULAR LOCATION: Nuclear (Potential).
CC -!- SIMILARITY: BELONGS TO THE SNF2/RAD54 HELICASE FAMILY. SNF2L
CC SUBFAMILY.
CC -----
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CC -----
DR EMBL; L27127; AAA19868.1; -;
DR EMBL; AE003821; AAF58479.1; -;
DR FlyBase; FBgn0011604; ISWI.
DR InterPro; IPR001410; DEAD.
DR InterPro; IPR001650; Helicase.C.
DR InterPro; IPR001005; MYD_DNA_bind.
DR Pfam; PF00271; helicase.C; 1.
DR Pfam; PF00176; SNF2_N; 1.
DR SMART; SM00487; SNF2_N; 1.
DR SMART; SM00487; DEXDC; 1.
DR SMART; SM00490; HELICc; 1.
DR SMART; SM00395; SANT; 2.
KW Nuclear protein; Helicase; ATP-binding.

FT NP_BIND 153 160 ATP (POTENTIAL).
FT SITE 256 259 DEAH BOX.
FT DOMAIN 978 981 POLY-LYS.
FT DOMAIN 1023 1027 POLY-LYS.
SQ SEQUENCE 1027 AA; 118873 MW; 008FC81AE15E071F CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1027;
Best Local Similarity 66.7%; Pred. No. 8.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 eagxxs 6
Db 658 EAGTSS 663

RESULT 28
PMAL_DICDI
ID PMAL_DICDI STANDARD; PRT; 1058 AA.
AC P54679;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Probable plasma membrane ATPase (EC 3.6.3.6) (Proton pump) (PAT2).
GN PATB.
OS Dictyostelium discoideum (Slime mold).
OC Eukaryota; Mycetoza; Dictyosteliida; Dictyostelium.
OX NCBI_TaxID=44689;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=AX3;
RX MEDLINE=98083743; PubMed=9421912;
RA Coukell M.B., Moniakis J., Cameron A.M.;
RT "The patB gene of Dictyostelium discoideum encodes a P-type H(+)-
RT ATPase isoform essential for growth and development under acidic
RT conditions.";
RL Microbiology 143:3877-3888(1997).
CC -!- FUNCTION: THE PLASMA MEMBRANE ATPASE IS A HYDROGEN ION PUMP. THE
CC PROTON GRADIENT IT GENERATES DRIVES THE ACTIVE TRANSPORT OF
CC NUTRIENTS BY H+ SYMPORT. THE RESULTING EXTERNAL ACIDIFICATION
CC AND/OR INTERNAL ALKINIZATION MAY MEDIATE GROWTH RESPONSES.
CC -!- CATALYTIC ACTIVITY: ATP + H(2O) + H(+)(IN) -> ADP + PHOSPHATE +
CC H(+)(OUT).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein.
CC -!- SIMILARITY: BELONGS TO THE CATION TRANSPORT ATPASES FAMILY
CC (E1-E2 ATPASES). SUBFAMILY I11A.
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CC -----
DR EMBL; X98286; CAA66931.1; -;
DR DictyDb; DD00061; patB.
DR InterPro; IPR004014; Cation_ATPase.
DR InterPro; IPR001757; E1-E2_ATPase.
DR InterPro; IPR000695; HATPase.
DR InterPro; IPR001454; Hydrolase.
DR Pfam; PF00690; Cation_ATPase_N; 1.
DR Pfam; PF00122; E1-E2_ATPase; 1.
DR Pfam; PF00702; Hydrolase; 1.
DR PRINTS; PR00119; CATATPASE.
DR PRINTS; PR00120; HATPASE.
DR PROSITE; PS00154; ATPase_E1_E2; 1.
KW Hydrolase; Hydrogen ion transport; Transmembrane; Phosphorylation;
KW Magnesium; ATP-binding.
FT DOMAIN 1 212 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 213 232 POTENTIAL.
FT DOMAIN 233 237 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 238 258 POTENTIAL.

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FT DOMAIN 259 387 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 388 407 POTENTIAL.
FT DOMAIN 388 417 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 408 417 POTENTIAL.
FT DOMAIN 426 447 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 448 473 POTENTIAL.
FT DOMAIN 784 805 POTENTIAL.
FT TRANSMEM 806 810 EXTRACELLULAR (POTENTIAL).
FT DOMAIN 811 833 POTENTIAL.
FT TRANSMEM 834 849 CYTOPLASMIC (POTENTIAL).
FT DOMAIN 850 870 POTENTIAL.
FT TRANSMEM 871 889 EXTRACELLULAR (POTENTIAL).
FT DOMAIN 890 910 POTENTIAL.
FT TRANSMEM 911 922 CYTOPLASMIC (POTENTIAL).
FT DOMAIN 923 943 POTENTIAL.
FT TRANSMEM 944 967 EXTRACELLULAR (POTENTIAL).
FT DOMAIN 968 988 POTENTIAL.
FT TRANSMEM 989 1058 CYTOPLASMIC (POTENTIAL).
FT MOD_RES 480 480 PHOSPHORYLATION (BY SIMILARITY).
FT METAL 728 732 MAGNESIUM (BY SIMILARITY).
FT METAL 732 732 MAGNESIUM (BY SIMILARITY).
FT DOMAIN 44 55 POLY-GLN.
FT DOMAIN 113 116 POLY-SER.
FT DOMAIN 246 249 POLY-LEU.
SQ SEQUENCE 1058 AA; 117373 MW; CB0E5AB9DEB9AF2 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1058;
Best Local Similarity 66.7%; Pred. No. 8.8e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxss 6
    ||| |
Db 110 EAGSSS 115

RESULT 29
ACAA_ARATH STANDARD; PRT; 1069 AA.
ID ACAA_ARATH
AC Q9S2R1; Q9M0D3;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Potential calcium-transporting ATPase 10, plasma membrane-type
DE (EC 3.6.3.8) (Ca2+-ATPase, isoform 10).
GS ACAL0 OR AT4G29900 OR F27B13.140.
ON Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosid II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
SEQUENCE FROM N.A.
RP STRAIN=CV. COLUMBIA;
RC MEDLINE=20083488; PubMed=10617198;
RX Mayer K.F.X., Schueller C., Wambutt R., Murphy G., Volckaert G.,
RA Pohl T., Duesterhoeft A., Stiekema W., Entian K.-D., Terryn N.,
RA Harris B., Ansoorge W., Brandt P., Grivell L.A., Rieger M.,
RA Weichselgartner M., de Simone V., Obermaier B., Mache R., Mueller M.,
RA Kreis M., Delsen M., Puigdomenech P., Watson M., Schmidtheini T.,
RA Reichert B., Portetelle D., Perez-Alonso M., Boutry M., Bancroft I.,
RA Vos P., Hoheisel J., Zimmermann W., Wedler H., Ridley P.,
RA Langham S.-A., McCullagh B., Bilham L., Robben J.,
RA Van der Schueren J., Grymonprez B., Chuang Y.-J., Vandenbussche F.,
RA Braekens M., Weltjens I., Voet M., Bastiaens I., Aert R., Defoor E.,
RA Weitzenegger T., Bothe G., Ramsperger U., Hilbert H., Braun M.,
RA Holzer E., Brandt A., Peters S., van Staveren M., Dirkse W.,
RA Moeljan P., Klein Lankhorst R., Rose M., Hauf J., Koetter P.,
RA Bernselser S., Hempel S., Feldpausch M., Lamberth S., van den Daele H.,
RA De Keyser A., Buysshaert C., Gielens J., Villarroel R., De Clercq R.,
RA Van Montagu M., Rogers J., Cronin A., Quail M., Bray-Allen S.,
RA Clark L., Doggett J., Hall S., Kay M., Lennard N., McLean K., Mayes R.,
RA Pettett A., Rajandream M.A., Lyne M., Benes V., Rechmann S.,
RA Borkova D., Bloeker H., Scharfe M., Grimm M., Loehnert T.-H.,

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RA Dose S., de Haan M., Maarse A.C., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fartmann B., Granderath K., Dauner D., Herzl A.,
RA Neumann S., Argiriou A., Vitale D., Liguori R., Piravandi E.,
RA Massenot O., Quigley F., Clabaud G., Muendlein A., Felber R.,
RA Schnabl S., Hiller R., Schmidt W., Lecharny A., Aubourg S.,
RA Chefor F., Cooke R., Berger C., Monfort A., Casacuberta E.,
RA Gibbons T., Weber N., Vandenbol M., Barges M., Terol J., Torres A.,
RA Perez-Perez A., Purnelle B., Bent E., Johnson S., Tacon D., Jesse T.,
RA Heijman D., Schwarz S., Scholler P., Heber S., Franks P., Bieleke C.,
RA Frishman D., Haase D., Lemcke K., Mewes H.-W., Stocker S.,
RA Zaccaria P., Bevan M., Willson R.K., de la Bastide M., Habermann K.,
RA Parnell L., Dedhia N., Gnoj L., Schutz K., Huang E., Spiegel L.,
RA Sekhon M., Murray J., Sheet P., Cordes M., Abu-Threideh J.,
RA Stoneking T., Kalicki J., Graves J., Harmon G., Edwards J.,
RA Latreille P., Courtney L., Cloud J., Abbott A., Scott K., Johnson D.,
RA Minx P., Bentley D., Fulton B., Miller N., Greco T., Kemp K.,
RA Kramer J., Fulton L., Mardis E., Dante M., Pepin K., Hillier L.,
RA Nelson J., Spieth J., Ryan E., Andrews S., Giesel C., Layman D.,
RA Du H., Ali J., Berghoff A., Jones K., Drone K., Cotton M., Joshi C.,
RA Antoniou B., Zidanic M., Strong C., Sun H., Lamar B., Yordan C.,
RA Ma P., Zhong J., Preston R., Vil D., Shekher M., Matero A., Shah R.,
RA Swaby I.K., O'Shaughnessy A., Rodriguez M., Hoffman J., Till S.,
RA Granat S., Shohdy N., Hasegawa A., Hameed A., Lodhi M., Johnson A.,
RA Chen E., Marra M., Martienssen R., McCombie W.R.;
RA 'Sequence and analysis of chromosome 4 of the plant Arabidopsis
RT thaliana.';
RT Nature 402:769-777(1999).
RL
CC -1- FUNCTION: THIS MAGNESIUM DEPENDENT ENZYME CATALYZES THE HYDROLYSIS
CC OF ATP COUPLED WITH THE TRANSLOCATION OF CALCIUM FROM THE CYTOSOL
CC INTO THE ENDOPLASMIC RETICULUM (BY SIMILARITY).
CC -1- CATALYTIC ACTIVITY: ATP + H(2)O + CA(2+)(CIS) = ADP + PHOSPHATE +
CC CA(2+)(TRANS).
CC -1- ENZYME REGULATION: ACTIVATED BY CALMODULIN (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- DOMAIN: THE N-TERMINUS CONTAINS AN AUTOINHIBITORY CALMODULIN-
CC BINDING DOMAIN, WHICH BINDS CALMODULIN IN A CALCIUM-DEPENDENT
CC FASHION (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE CATION TRANSPORT ATPASES FAMILY
CC (E1-E2 ATPASES). SUBFAMILY IIB.
CC -1- CAUTION: THE SEQUENCE CAB43665 DIFFERS FROM THAT SHOWN DUE TO
CC WRONG EXON BOUNDARIES PREDICTED FROM THE GENOMIC SEQUENCE.
CC -----
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AL161575; CAB79748.1;
CC EMBL; AL050352; CAB43665.1; ALT_SEQ.
CC HSSP; P04191; LEUL.
CC InterPro; IPR004014; Cation_ATPase.
CC InterPro; IPR001757; E1-E2_ATPase.
CC InterPro; IPR001454; Hydrolase.
CC Pfam; PF00689; Cation_ATPase_C; 1.
CC Pfam; PF00690; Cation_ATPase_N; 1.
CC Pfam; PF00122; E1-E2_ATPase; 1.
CC Pfam; PF00702; Hydrolase; 1.
CC PRINTS; PR00119; CATATPASE.
CC PROSITE; PS00134; ATPASE_E1_E2; 1.
CC Hydrolase; Calcium transport; Transmembrane; Phosphorylation;
CC ATP-binding; Metal-binding; Magnesium; Calmodulin-binding;
CC Multigene family.
CC DOMAIN 1 180 CYTOPLASMIC (POTENTIAL).
CC TRANSMEM 181 201 POTENTIAL.
CC DOMAIN 202 219 LUMENAL (POTENTIAL).
CC TRANSMEM 220 240 POTENTIAL.
CC DOMAIN 241 369 CYTOPLASMIC (POTENTIAL).
CC TRANSMEM 370 389 POTENTIAL.
CC DOMAIN 390 426 LUMENAL (POTENTIAL).
CC TRANSMEM 427 444 POTENTIAL.

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FT DOMAIN 445 844 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 845 863 POTENTIAL.
FT DOMAIN 864 874 LUMENAL (POTENTIAL).
FT TRANSMEM 875 895 POTENTIAL.
FT DOMAIN 896 915 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 916 938 POTENTIAL.
FT DOMAIN 939 950 LUMENAL (POTENTIAL).
FT TRANSMEM 952 973 POTENTIAL.
FT DOMAIN 974 991 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 992 1013 POTENTIAL.
FT DOMAIN 1014 1023 LUMENAL (POTENTIAL).
FT TRANSMEM 1024 1045 POTENTIAL.
FT DOMAIN 1046 1069 CYTOPLASMIC (POTENTIAL).
FT MOD_RES 482 482 CALMODULIN-BINDING (BY SIMILARITY).
FT METAL 789 789 PHOSPHORYLATION (BY SIMILARITY).
FT METAL 793 793 MAGNESIUM (BY SIMILARITY).
SQ SEQUENCE 1069 AA; 116858 MW; 48CE4A4B218E2656 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1069;
Best Local Similarity 66.7%; Pred. No. 8.9e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
DB 17 EAGTSS 22

RESULT 30
CNA_STRAU STANDARD; PRT: 1183 AA.
AC Q53654;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Collagen adhesin precursor.
GN CNA.
OS Staphylococcus aureus.
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Staphylococcus.
OX NCBI_TaxID=1280;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=FDA 574;
RX MEDLINE=92165839; PubMed=1311320;
RA Patti J.M., Jonsson H., Guss B., Switalski L.M., Wiberg K.,
RA Lindberg M., Hoeoek M.;
RT "Molecular characterization and expression of a gene encoding a
RT Staphylococcus aureus collagen adhesin.";
RL J. Biol. Chem. 267:4766-4772(1992).
RN [2]
RP ERRATUM.
RA Patti J.M., Jonsson H., Guss B., Switalski L.M., Wiberg K.,
RA Lindberg M., Hoeoek M.;
RL J. Biol. Chem. 269:11672-11672(1994).
RN [3]
RP COLLAGEN-BINDING DOMAIN.
RC STRAIN=FDA 574;
RX MEDLINE=94032261; PubMed=8218209;
RA Patti J.M., Boles J.O., Hoeoek M.;
RT "Identification and biochemical characterization of the ligand
RT binding domain of the collagen adhesin from Staphylococcus aureus.";
RL Biochemistry 32:11428-11435(1993).
RN [4]
RP X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS) OF 169-318.
RX MEDLINE=97475225; PubMed=9334749;
RA Symerksy J., Patti J.M., Carlson M., House-Pompeo K., Teale M.,
RA Moore D., Jin L., Schneider A., Delucas L.J., Hoeoek M.,
RA Narayana S.V.L.;
RT "Structure of the collagen-binding domain from a Staphylococcus
RT aureus adhesin.";
RL Nat. Struct. Biol. 4:833-838(1997).

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CC -!- FUNCTION: MEDIATES ATTACHMENT OF STAPHYLOCOCCAL CELLS TO
CC COLLAGEN-CONTAINING SUBSTRATA.
CC -!- SUBCELLULAR LOCATION: Type I membrane protein. Cell wall.
CC -!- SIMILARITY: TO OTHER STREPTOCOCCAL AND STAPHYLOCOCCAL PROTEINS
CC IN THE REGION OF THE MEMBRANE ANCHOR.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: M81736; AAA20874.1; -.
CC PDB: 1AMX; 24-JUN-98
CC InterPro: IPR001899; Gram_pos_anchor.
CC PROSITE: PS00343; GRAM_POS_ANCHORING; FALSE_NEG.
CC Signal; Repeat; Transmembrane; Cell wall; 3D-structure.
CC SIGNAL 1 29 POTENTIAL.
CC CHAIN 30 1183 COLLAGEN ADHESIN.
CC DOMAIN 30 1157 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 1158 1177 MEMBRANE ANCHOR (POTENTIAL).
CC DOMAIN 1178 1183 CYTOPLASMIC (POTENTIAL).
CC DOMAIN 151 318 COLLAGEN-BINDING.
CC DOMAIN 533 1093 3 X 187 AA APPROXIMATE TANDEM REPEATS.
CC DOMAIN 1093 1157 LYS/PRO-RICH (CELL WALL-SPANNING).
CC DOMAIN 1151 1156 CONSERVED IN GRAM-POSITIVE COCCI SURFACE
CC PROTEINS.
CC REPEAT 533 719 B1.
CC REPEAT 720 906 B2.
CC REPEAT 907 1093 B3.
CC SEQUENCE 1183 AA; 133066 MW; B6A1CC072E575D76 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1183;
Best Local Similarity 66.7%; Pred. No. 9.8e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
DB 166 EAGTSS 171

RESULT 31
B3A2_MOUSE STANDARD; PRT: 1237 AA.
ID B3A2_MOUSE Q9ES13; Q9ES12; Q9ES11; Q9ES10; Q9ES09;
AC F13808; Q9ES13; Q9ES12; Q9ES11; Q9ES10; Q9ES09;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Anion exchange protein 2 (Non-erythroid band 3-like protein) (B3RP).
GN SLC4A2 OR AE2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM A).
RX MEDLINE=89034212; PubMed=3182834;
RA Alper S.L., Kopito R.R., Libresco S.M., Lodish H.F.;
RT "Cloning and characterization of a murine band 3-related cDNA from
RT kidney and from a lymphoid cell line.";
RL J. Biol. Chem. 263:17092-17099(1988).
RN [2]
RP SEQUENCE FROM N.A., ALTERNATIVE SPLICING, AND TISSUE SPECIFICITY.
RX MEDLINE=20462926; PubMed=11006093;
RA Lecanda J., Urtasun R., Medina J.F.;
RT "Molecular cloning and genomic organization of the mouse AE2 anion
RT exchanger gene.";
RL Biochem. Biophys. Res. Commun. 276:117-124(2000).
CC -!- FUNCTION: PLASMA MEMBRANE ANION EXCHANGE PROTEIN OF WIDE
CC DISTRIBUTION.

```

CC -!- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -!- ALTERNATIVE PRODUCTS: 5 isoforms; a (shown here), b1, b2, c1 and
 CC c2; are produced by alternative splicing.
 CC -!- TISSUE SPECIFICITY: Isoform a is widely expressed at similar
 CC levels in all tissues examined. Isoforms b1 and b2 are
 CC predominantly expressed in stomach although they are also detected
 CC at lower levels in other tissues. Isoform c1 is stomach-specific.
 CC Isoform c2 is expressed at slightly higher levels in lung and
 CC stomach than in other tissues.
 CC -!- SIMILARITY: BELONGS TO THE ANION EXCHANGER FAMILY.
 CC -----
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DR EMBL; J04036; AAA65505.1; -.
 DR EMBL; AF255774; AAG23154.1; -.
 DR EMBL; AF255774; AAG23155.1; -.
 DR EMBL; AF255774; AAG23156.1; -.
 DR EMBL; AF255774; AAG23157.1; -.
 DR EMBL; AF255774; AAG23158.1; -.
 DR EMBL; AF255774; AAG23159.1; -.
 DR PIR; A31789; A31789.
 DR HSP; P02730; 1BTR.
 DR MGD; MGI:109351; Slc4a2.
 DR InterPro: IPR001717; Anion_exchanger.
 DR InterPro: IPR003020; HCO3_cotransp.
 DR Pfam; PF00955; HCO3_cotransp; 1.
 DR PRINTS; PR01231; HCO3TRNSPOT.
 DR PROSITE; PS00219; ANION_EXCHANGER_1; 1.
 DR PROSITE; PS00220; ANION_EXCHANGER_2; 1.
 KW Transmembrane; Glycoprotein; Anion exchange; Lipoprotein; Palmitate;
 KW Alternative splicing.
 FT DOMAIN 1 703 CYTOPLASMIC (POTENTIAL).
 FT DOMAIN 704 1237 MEMBRANE (ANION EXCHANGE).
 FT TRANSMEM 704 727 POTENTIAL.
 FT TRANSMEM 733 770 POTENTIAL.
 FT TRANSMEM 790 812 POTENTIAL.
 FT TRANSMEM 822 843 POTENTIAL.
 FT DOMAIN 844 896 EXPLOSMIC LOOP (POTENTIAL).
 FT TRANSMEM 897 914 POTENTIAL.
 FT DOMAIN 915 929 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 930 950 POTENTIAL.
 FT TRANSMEM 984 1006 POTENTIAL.
 FT TRANSMEM 1032 1053 POTENTIAL.
 FT TRANSMEM 1087 1132 POTENTIAL.
 FT TRANSMEM 1159 1195 POTENTIAL.
 FT DOMAIN 5 316 PRO-RICH.
 FT DOMAIN 73 87 HIS-RICH.
 FT DOMAIN 861 865 POLY-SER.
 FT CARBOHYD 855 855 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 866 866 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 878 878 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT LIPID 1169 1169 PALMITATE (BY SIMILARITY).
 FT VARSPPLIC 1 17 MSSAPRRPSSAGDSLHT -> MDPLRPQ (IN ISOFORM B2).
 FT VARSPPLIC 1 17 MSSAPRRPSSAGDSLHT -> MTQ (IN ISOFORM B1).
 FT VARSPPLIC 1 166 MISSING (IN ISOFORM C2).
 FT VARSPPLIC 1 193 MISSING (IN ISOFORM C1).
 FT VARSPPLIC 167 198 ERTSPSPPTQPHQEAAPRASKGAQTG -> MPFAQEWKSG
 FT FT GLREAVFGAGGSGVCR (IN ISOFORM C2).
 FT FT A -> G (IN REF. 2).
 FT CONFLICT 205 205
 FT SEQUENCE 1237 AA; 136813 MW; 1A0782C0071782EE CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1237;
 Best Local Similarity 66.7%; Pred. No. 1e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 eagxxs 6

Db 858 EAGSSS 863
 RESULT 32
 MYO6_HUMAN STANDARD; PRT; 1262 AA.
 ID Q9UM54;
 AC Q9UM54;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 01-MAR-2002 (Rel. 41, Last annotation update)
 DE Myosin VI.
 GN MYO6.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OC NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97402203; PubMed=9259267;
 RA Avraham K.B., Hasson T., Sobel T., Balsara B., Testa J.R.,
 RA Skvorak A.B., Morton C.C., Copeland N.G., Jenkins N.A.;
 RT "Characterization of unconventional MYO6, the human homologue of the
 RT gene responsible for deafness in Snell's waltzer mice.";
 RL Hum. Mol. Genet. 6:1225-1231(1997).
 RN [2]
 RP REVISIONS.
 RA Avraham K.B.;
 RL Submitted (JUL-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP FUNCTION.
 RX MEDLINE=99447046; PubMed=10519557;
 RA Wells A.L., Lin A.W., Chen L.-Q., Safer D., Cain S.M., Hasson T.,
 RA Carragher B.O., Milligan R.A., Sweeney H.L.;
 RT "Myosin VI is an actin-based motor that moves backwards.";
 RL Nature 401:505-508(1999).
 RN [4]
 RP VARIANT DFNA22 TYR-442.
 RX MEDLINE=21375673; PubMed=11468689;
 RA Melchionda S., Ahituv N., Biscaglia L., Sobel T., Glaser F.,
 RA Rabionet R., Arbones M.L., Notarangelo A., Di Iorio E., Catella M.,
 RA Zelante L., Estivill X., Avraham K.B., Gasparini P.;
 RT "MYO6, the human homologue of the gene responsible for deafness in
 RT Snell's waltzer mice, is mutated in autosomal dominant nonsyndromic
 RT hearing loss.";
 RL Am. J. Hum. Genet. 69:635-640(2001).
 CC -!- FUNCTION: RECESSIVE ACTIN-BASED MOTOR. REQUIRED FOR STRUCTURAL
 CC INTEGRITY OF INNER EAR HAIR CELLS (BY SIMILARITY).
 CC -!- DISEASE: Defects in MYO6 are the cause of an autosomal dominant
 CC form of nonsyndromic sensorineural deafness (DFNA22). The deafness
 CC is progressive and postlingual, with onset during childhood (8 to
 CC 10 years of age at onset of symptoms; 6 to 8 years of age at onset
 CC of first audiometric abnormalities). By the age of approximately
 CC 50 years, affected individuals invariably have profound
 CC sensorineural deafness.
 CC -!- SIMILARITY: CONTAINS 1 MYOSIN-LIKE GLOBULAR HEAD DOMAIN.
 CC -!- SIMILARITY: CONTAINS 1 IQ DOMAIN.
 CC -----
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 CC -----
 CC EMBL; U90236; AAC51654.2; -.
 DR HSP; P08799; ILVK.
 DR MIM; 600970; -.
 DR MIM; 606346; -.
 DR InterPro: IPR000048; IQ.
 DR InterPro: IPR001609; myosin_head.
 DR Pfam; PF00612; IQ; 1.

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DR Pfam: PF00063; myosin_head; 1.
DR PRINTS: PR00193; MYOSINHEAVY.
DR ProDom: PD000355; myosin_head; 1.
DR SMART: SM00015; IQ; 1.
DR SMART: SM00242; MYSC; 1.
DR PROSITE: PS00096; IQ; FALSE_NEG.
KW Myosin; ATP-binding; Calmodulin-binding; Actin-binding;
KW Coiled coil; Disease mutation; Deafness.
FT DOMAIN 1 759 MYOSIN HEAD-LIKE.
FT DOMAIN 814 834 IQ.
FT DOMAIN 848 1030 COILED COIL (POTENTIAL).
FT NP_BIND 151 158 ATP (POTENTIAL).
FT DOMAIN 665 672 ACTIN-BINDING (POTENTIAL).
FT VARIANT 442 442 C -> Y (IN DFNA22).
FT /FTID=VAR_012110.
SQ SEQUENCE 1262 AA; 146047 MW; CF1FA35796FC1C60 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1262;
Best Local Similarity 66.7%; Pred. No. 1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxys 6
Db 354 EAGSTS 359.

RESULT 33
DYNA_DROME STANDARD; PRT; 1265 AA.
AC F13496; Q9VUAI;
DT 01-JAN-1990 (Rel. 13, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE 150 kDa dynein-associated polypeptide (DP-150) (Glued
DE protein).
DE GL OR C69206.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
[1]
[2]
SEQUENCE FROM N.A.
RN STRAIN=OREGON-R, AND CANTON-S;
RX MEDLINE=87317680; PubMed=2819881;
RA Swatcoop A., Swatcoop M., Garen A.;
RT "Sequence analysis of the complete cDNA and encoded polypeptide for
RT the Glued gene of Drosophila melanogaster.";
RL Proc. Natl. Acad. Sci. U.S.A. 84:6501-6505(1987).
RN [2]
SEQUENCE FROM N.A.
RN STRAIN=BERKELEY;
RX MEDLINE=20196006; PubMed=10731132;
RA Adams M.D., Celnik S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Ananthanades P.G., Scher S.E., Li P.W., Hoskins R.A., Galie R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
RA Bailew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
RA Borokova D., Botchan M.R., Bouck J., Brokstein P., Brotter P.,
RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
RA Foster C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Hartis N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,

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RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wassarman D.A., Weinstein G.M., Weissenbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2185-2195(2000).
CC -!- FUNCTION: REQUIRED FOR THE CYTOPLASMIC DYNEIN-DRIVEN RETROGRADE
CC MOVEMENT OF VESICLES AND ORGANELLES ALONG MICROTUBULES. DYNEIN-
CC DYNACTIN INTERACTION IS A KEY COMPONENT OF THE MECHANISM OF AXONAL
CC TRANSPORT OF VESICLES AND ORGANELLES.
CC -!- SUBUNIT: LARGE MACROMOLECULAR COMPLEX OF AT LEAST 10 COMPONENTS.
CC P150(GLUED) BINDS DIRECTLY TO MICROTUBULES AND TO CYTOPLASMIC
CC DYNEIN.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: BELONGS THE DYNACTIN 150 KDA SUBUNIT FAMILY.
CC -!- SIMILARITY: CONTAINS 1 CAP-GLY DOMAIN.
CC -!- CAUTION: REF.1 SEQUENCE DIFFERS FROM THAT SHOWN DUE TO
CC FRAMESHIFTS AT POSITIONS 32; 174 TO 220; 648 TO 672 AND 1208.
CC -----
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CC -----
DR EMBL: J02932; -; NOT_ANNOTATED_CDS.
DR EMBL: AE003536; AAF49788.1; -.
DR F1R; A28313; A28313.
DR FlyBase: FBgn0001108; GL.
DR InterPro: IPR000938; CAP-Gly.
DR Pfam: PF01302; CAP_GLY; 1.
DR PROSITE: PS00845; CAP_GLY_1; 1.
DR PROSITE: PS50245; CAP_GLY_2; 1.
KW Motor protein; Microtubules; Dynein; Coiled coil; Cytoskeleton.
FT DOMAIN 27 69 CAP-GLY.
FT DOMAIN 105 138 SER-RICH.
FT DOMAIN 213 570 COILED COIL (POTENTIAL).
FT DOMAIN 812 836 COILED COIL (POTENTIAL).
FT DOMAIN 967 1084 COILED COIL (POTENTIAL).
FT DOMAIN 1128 1160 COILED COIL (POTENTIAL).
FT CONFLICT 708 708 D -> A (IN REF. 1).
FT CONFLICT 875 875 L -> V (IN REF. 1).
FT CONFLICT 888 888 A -> R (IN REF. 1).
FT CONFLICT 1043 1043 S -> C (IN REF. 1).
SQ SEQUENCE 1265 AA; 141217 MW; 2038A200282B2755 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1265;
Best Local Similarity 66.7%; Pred. No. 1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxys 6
Db 802 EAGATS 807

RESULT 34
MY06_MOUSE

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ID MYO6_MOUSE STANDARD; PRT; 1265 AA.
AC Q64331;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Myosin VI.
GN MYO6 OR SV.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RX MEDLINE=96083582; PubMed=7493015;
RA Avraham K.B., Hasson T., Steel K.P., Kingsley D.M., Russell L.B.,
RA Moosker M.S., Copeland N.G., Jenkins N.A.;
RT "The mouse Snell's waltzer deafness gene encodes an unconventional
RT myosin required for structural integrity of inner ear hair cells.";
RL Nat. Genet. 11:369-375(1995).
CC -1- FUNCTION: RECESSIVE ACTIN-BASED MOTOR. REQUIRED FOR STRUCTURAL
CC INTEGRITY OF INNER EAR HAIR CELLS (BY SIMILARITY).
CC -1- TISSUE SPECIFICITY: WIDELY EXPRESSED. WITHIN THE COCHLEA,
CC EXPRESSED SPECIFICALLY WITHIN THE SENSORY HAIR CELLS.
CC -1- DISEASE: DEFECTS IN MYO6 ARE THE CAUSE OF SNELL'S WALTZER, A
CC CONDITION CHARACTERIZED BY CIRCLING, HEAD-TOSSING, DEAFNESS AND
CC HYPERACTIVITY.
CC -1- SIMILARITY: CONTAINS 1 MYOSIN-LIKE GLOBULAR HEAD DOMAIN.
CC -1- SIMILARITY: CONTAINS 1 IQ DOMAIN.
CC
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CC
CC EMBL; U49739; AAB00194.1; -.
CC HSP; P08796; ILVK.
CC MGD; MG1:104785; Myo6.
CC InterPro; IPR000048; IQ.
CC InterPro; IPR001609; myosin_head.
CC Pfam; PF00612; IQ; 1.
CC Pfam; PF00063; myosin_head; 4.
CC PRINTS; PR00193; MYOSINHEAVY.
CC ProDom; PD000355; myosin_head; 1.
CC SMART; SM00015; IQ; 1.
CC SMART; SM00242; MYSC; 1.
CC PROSITE; PS50096; IQ; FALSE_NEG.
KW Myosin; ATP-binding; Calmodulin-binding; Actin-binding;
KW Coiled coil; Disease mutation; Deafness.
FT DOMAIN 1 762 MYOSIN HEAD-LIKE.
FT DOMAIN 817 837 IQ.
FT DOMAIN 849 1014 COILED COIL (POTENTIAL).
FT NP_BIND 151 158 ATP (POTENTIAL).
FT DOMAIN 668 1265 ACTIN-BINDING (POTENTIAL).
FT VARIANT 766 1265 MISSING (IN SNELL'S WALTZER).
SQ SEQUENCE 1265 AA; 14640 MW; 4F51ABC72463148C CRC64;

Query Match 90.58; Score 19; DB 1; Length 1265;
Best Local Similarity 66.78; Pred. No. 1e-03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
DB 355 EAGSTS 360
||| |
RP SEQUENCE FROM N.A. (ISOFORM 2).
RC STRAIN=C57BL/6N;
RA Wang Z., Goldstein A., Neufeld E.J., Scheuermann R.H., Tucker P.W.;
RT "Repression of immunoglobulin heavy chain intronic enhancer
RT through nuclear matrix attachment sites: Cux/CDP homeoprotein is a
RT component of NF-muNR repressor.";
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE OF 64-1395 FROM N.A. (ISOFORM 1).

ID Y232_HUMAN STANDARD; PRT; 1278 AA.
AC Q92628;
DT 01-NOV-1997 (Rel. 35, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hypothetical protein KIAA0232 (Fragment).
GN KIAA0232.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Nagase T., Seki N., Ishikawa K.-I., Ohira M., Kawarabayasi Y.,
RA Ohara O., Tanaka A., Kotani H., Miyajima N., Nomura N.;
RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE OF 271-1278 FROM N.A.
RC TISSUE=Bone marrow;
RX MEDLINE=97191544; PubMed=9039502;
RA Nagase T., Seki N., Ishikawa K.-I., Ohira M., Kawarabayasi Y.,
RA Ohara O., Tanaka A., Kotani H., Miyajima N., Nomura N.;
RT "Prediction of the coding sequences of unidentified human genes. VI.
RT The coding sequences of 80 new genes (KIAA0201-KIAA0280) deduced by
RT analysis of cDNA clones from cell line KG-1 and brain.";
RL DNA Res. 3:321-329(1996).
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; D86985; BAA13221.2; -.
CC KW Hypothetical protein.
CC NON_TER 1
FT NON_TER 1
SQ SEQUENCE 1278 AA; 141663 MW; 2FCFC8837AF8134D CRC64;

Query Match 90.58; Score 19; DB 1; Length 1278;
Best Local Similarity 66.78; Pred. No. 1e-03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
DB 177 EAGSSS 182
||| |
RP SEQUENCE FROM N.A. (ISOFORM 2).
RC STRAIN=C57BL/6N;
RA Wang Z., Goldstein A., Neufeld E.J., Scheuermann R.H., Tucker P.W.;
RT "Repression of immunoglobulin heavy chain intronic enhancer
RT through nuclear matrix attachment sites: Cux/CDP homeoprotein is a
RT component of NF-muNR repressor.";
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE OF 64-1395 FROM N.A. (ISOFORM 1).

ID Y232_HUMAN STANDARD; PRT; 1278 AA.
AC Q92628;
DT 01-NOV-1997 (Rel. 35, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hypothetical protein KIAA0232 (Fragment).
GN KIAA0232.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Nagase T., Seki N., Ishikawa K.-I., Ohira M., Kawarabayasi Y.,
RA Ohara O., Tanaka A., Kotani H., Miyajima N., Nomura N.;
RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE OF 271-1278 FROM N.A.
RC TISSUE=Bone marrow;
RX MEDLINE=97191544; PubMed=9039502;
RA Nagase T., Seki N., Ishikawa K.-I., Ohira M., Kawarabayasi Y.,
RA Ohara O., Tanaka A., Kotani H., Miyajima N., Nomura N.;
RT "Prediction of the coding sequences of unidentified human genes. VI.
RT The coding sequences of 80 new genes (KIAA0201-KIAA0280) deduced by
RT analysis of cDNA clones from cell line KG-1 and brain.";
RL DNA Res. 3:321-329(1996).
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; D86985; BAA13221.2; -.
CC KW Hypothetical protein.
CC NON_TER 1
FT NON_TER 1
SQ SEQUENCE 1278 AA; 141663 MW; 2FCFC8837AF8134D CRC64;
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RC STRAIN-A/J, AND BALB/C; TISSUE=Brain;
RX MEDLINE=94244481; PubMed=7910552;
RA Valarche I., Tissier-Seta J.P., Hirsch M.R., Martinez S., Goridis C.,
RA Brunet J.F.;
RT "The mouse homeodomain protein Phox2 regulates Ncam promoter activity
in concert with Cux/CDP and is a putative determinant of
neurotransmitter phenotype.";
RL Development 119:881-896(1993).
RN [3]
RP SEQUENCE OF 642-1395 FROM N.A.
RX MEDLINE=96437626; PubMed=8840273;
RA den Heuvel G.B., Bodmer K., McConnell K.R., Nagami G.T., Igarashi P.;
RT "Expression of a cut-related homeobox gene in developing and
polycystic mouse kidney.";
RL Kidney Int. 50:453-461(1996).
RN [4]
RP SEQUENCE OF 936-1395 FROM N.A.
RC TISSUE=Testis;
RA Quaglin S.E., Igarashi P.;
RT "A unique variant of a homeobox gene related to Drosophila cut is
expressed in mouse testis.";
RL Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: PROBABLY HAS A BROAD ROLE IN MAMMALIAN DEVELOPMENT AS A
REPRESSOR OF DEVELOPMENTALLY REGULATED GENE EXPRESSION. MAY ACT BY
PREVENTING BINDING OF POSITIVELY-ACTIVATING COAT FACTORS TO
PROMOTERS (BY SIMILARITY). COMPONENT OF NF-MUNR REPRESSOR; BINDS
TO THE MARS (5' AND 3') OF THE IMMUNOGLOBULIN HEAVY CHAIN
ENHANCER.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS; 1 (SHOWN HERE) AND 2; ARE
PRODUCED BY ALTERNATIVE SPLICING.
CC -1- SIMILARITY: CONTAINS 3 CUT DOMAINS.
CC -1- SIMILARITY: BELONGS TO THE CUT FAMILY OF HOMEBOX PROTEINS.
CC -----
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CC -----
DR EMBL; AF004225; AAD12485.1; -;
DR EMBL; X75013; CAA52922.1; -;
DR EMBL; U46683; AAC52775.1; -;
DR EMBL; U46684; AAB41146.1; -;
DR HSSP; P10037; 1AU7;
DR MGD; MGI:88568; Cut11.
DR InterPro; IPR003350; CUT.
DR InterPro; IPR001356; Homeobox.
DR Pfam; PF02376; CUT; 3.
DR Pfam; PF00046; homeobox; 2.
DR SMART; SM00389; HOX; 1.
DR PROSITE; PS00027; HOMEBOX_1; 1.
DR PROSITE; PS00071; HOMEBOX_2; 1.
DR Transcription regulation; Homeobox; DNA-binding;
KW Developmental protein; Nuclear protein; Repeat; Repressor;
KW Coiled coil; Alternative splicing.
FT NON_TER 1
FT DOMAIN 1 243 COILED COIL (POTENTIAL).
FT DNA_BIND 420 507 CUT 1.
FT DOMAIN 547 603 COILED COIL (POTENTIAL).
FT DNA_BIND 809 896 CUT 2.
FT DNA_BIND 992 1079 CUT 3.
FT DNA_BIND 1119 1178 HOMEBOX.
FT VARSPIC 1287 1388 MISSING (IN ISOFORM 2).
FT CONFLICT 1360 1360 G -> A (IN REF. 2).
FT CONFLICT 1365 1365 P -> L (IN REF. 1).
SQ SEQUENCE 1395 AA; 151802 MW; D062CC227D7A163E CRC64;

Query Match

Best Local Similarity 90.5%; Score 19; DB 1; Length 1395;

96.7%; Pred. No. 1.2e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 eaqxss 6
Db 377 EAGSTS 382
RESULT 37
MUKB_ECOLI STANDARD; PRT; 1486 AA.
ID MUKB_ECOLI
AC P22323; P77164; Q47398;
DT 01-AUG-1991 (Rel. 19, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Cell division protein mukB.
GN MUKB OR B0924.
OS Escherichia coli.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
OX NCBI_TaxID=562;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-K12 / W3110;
RX MEDLINE=91114703; PubMed=1989883;
RA Niki H., Jaffe A., Inamura R., Ogura T., Hiraga S.;
RT "The new gene mukB codes for a 177 kd protein with coiled-coil
domains involved in chromosome partitioning of E. coli.";
RL EMBO J. 10:183-193(1991).
RN [2]
RP SEQUENCE FROM N.A., AND MUTANTS MUKB33 AND MUKB106.
RX MEDLINE=95080615; PubMed=7988894;
RA Yamanaka K., Mitani T., Feng J., Ogura T., Niki H., Hiraga S.;
RT "Two mutant alleles of mukB, a gene essential for chromosome
partition in Escherichia coli.";
RL FEMS Microbiol. Lett. 123:27-31(1994).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;
RA Blattner F.R., Plunkett G. III, Bloch C.A., Perna N.T., Burland V.,
RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
RA Mau B., Shao Y.;
RT "The complete genome sequence of Escherichia coli K-12.";
RL Science 277:1453-1474(1997).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN-K12;
RX MEDLINE=97061202; PubMed=8905232;
RA Oshima T., Aiba H., Baba T., Fujita K., Hayashi K., Honjo A.,
RA Ikemoto K., Inada T., Itoh T., Kajihara M., Kanai K., Kashimoto K.,
RA Kimura S., Kitagawa M., Makino K., Masuda S., Miki T., Mizobuchi K.,
RA Mori H., Motomura K., Nakamura Y., Nishimoto H., Nishio Y., Saito N.,
RA Sampei G., Seki Y., Tagami H., Takemoto K., Wada C., Yamamoto Y.,
RA Yano M., Horiuchi T.;
RT "A 718-kb DNA sequence of the Escherichia coli K-12 genome
corresponding to the 12.7-28.0 min region on the linkage map.";
RL DNA Res. 3:137-155(1996).
RN [5]
RP SEQUENCE OF 1-44 FROM N.A.
RC STRAIN-K12 / W3110;
RX MEDLINE=94232180; PubMed=7513784;
RA Feng J., Yamanaka K., Niki H., Ogura T., Hiraga S.;
RT "New killing system controlled by two genes located immediately
upstream of the mukB gene in Escherichia coli.";
RL Mol. Gen. Genet. 243:136-147(1994).
RN [6]
RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 1-227.
RX MEDLINE=20015369; PubMed=10545328;
RA van den Ent F., Lockhart A., Kendrick-Jones J., Loewe J.;
RT "Crystal structure of the N-terminal domain of MukB: a protein
involved in chromosome partitioning.";
RL Structure 7:1181-1187(1999).

CC -1- FUNCTION: ESSENTIAL FOR CHROMOSOME PARTITIONING. IMPLICATED IN
CC ATP-DEPENDENT CHROMOSOME PARTITIONING DURING CELL DIVISION.
CC -1- SIMILARITY: CONTAINS A COILED COIL MYOSIN-LIKE STRUCTURE.
CC -----
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DR EMBL; X57550; CAA0776.1; -;
DR EMBL; D31701; BAA06510.1; -;
DR EMBL; AE000194; AAC74010.1; -;
DR EMBL; D90730; BAA35670.1; -;
DR EMBL; D26440; BAA05459.1; -;
DR PIR; JH0228; JH0228;
DR PDB; 1OHL; 10-NOV-99.
DR EcoGene; EG10618; mukB.
KW ATP-binding; Coiled coil; 3D-structure; Complete proteome.
FT DOMAIN 331 665 COILED COIL (POTENTIAL).
FT DOMAIN 784 1116 COILED COIL (POTENTIAL).
FT DOMAIN 1209 1265 COILED COIL (POTENTIAL).
FT NP_BIND 34 41 ATP (BY SIMILARITY).
FT VARIANT 33 33 S -> F (IN MUKB106).
FT VARIANT 1201 1201 D -> N (IN MUKB33).
FT CONFLICT 266 266 A -> R (IN REF. 1 AND 2).
FT CONFLICT 318 319 EH -> DD (IN REF. 1).
FT CONFLICT 1134 1134 H -> D (IN REF. 1).
FT CONFLICT 1174 1175 SE -> VQ (IN REF. 1).
FT CONFLICT 1276 1277 MISSING (IN REF. 1 AND 2).
FT CONFLICT 1357 1380 WLRKESGALSTGAIGTGMISLYM -> CCAQSLVHCRPVVR
FT FT WRYRYVDSGV (IN REF. 1).
FT FT SRLRGKDIPSKRTLLFLDEAARLDARSATLFLCERLQWQ
FT FT LITAAENISPERKTYKLVKRVQNTSEHVHVVLGRFAPO
FT FT LPETLPGTDEAPQAS -> SAACAVKISLLAACSSMKOR
FT FT DWMLVSLPCLNCSVCKNSQSSQRRKISARRKAPPINWCV
FT FT KSSRIPTNFMSSACEDLRNLSKRFQELTKRLRLRVKIKQO
FT FT CRLLFFKRLFCFKVAHYGALFFKLLIYRLCKNVRRLYTE
FT FT DKPDE (IN REF. 1).
SQ SEQUENCE 1486 AA; 170229 MW; 38C7874BEB78D6D6 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1486;
Best Local Similarity 66.7%; Pred. No. 1.2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
| | | | |
Db 65 EAGATS 70

RESULT 38
CUTL1_HUMAN
ID CUTL1_HUMAN STANDARD; PRT; 1505 AA.
AC P39880; Q9UEV5;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE CCAAT displacement protein (CDP) (Cut-like 1).
GN CUTL1.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Umbilical vein;
RX MEDLINE=93250973; PubMed=1301999;
RA Neufeld E.J., Skalnik D.G., Lievens P.M.-J., Orkin S.H.;
RT "Human CCAAT displacement protein is homologous to the Drosophila
homeoprotein, cut.";

RL Nat. Genet. 1:50-55(1992).
RN [2]
RP SEQUENCE OF 48-224 FROM N.A.
RX MEDLINE=99018118; PubMed=9799793;
RA Gloeckner G., Scherer S., Schattevov R., Boright A., Weber J.,
RA Tsui L.-C., Rosenthal A.,
RT "large-scale sequencing of two regions in human chromosome 7q22:
RT analysis of 650 kb of genomic sequence around the EPO and CUTL1 loci
RT reveals 17 genes.";
RL Genome Res. 8:1060-1073(1998).
CC -1- FUNCTION: PROBABLY HAS A BROAD ROLE IN MAMMALIAN DEVELOPMENT AS A
CC REPRESSOR OF DEVELOPMENTALLY REGULATED GENE EXPRESSION. MAY ACT BY
CC PREVENTING BINDING OF POSITIVELY-ACTING CCAAT FACTORS TO
CC PROMOTERS.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- ALTERNATIVE PRODUCTS: AT LEAST 2 ISOFORMS; 1 (SHOWN HERE) AND 2;
CC ARE PRODUCED BY ALTERNATIVE SPLICING.
CC -1- DOMAIN: ASN AT POSITION 47 OF THE HOMEBOX MAY PARTICIPATE IN
CC REGULATING DNA-BINDING ACTIVITY BY PROMOTING HOMO- AND
CC HETERODIMERIZATION.
CC -1- SIMILARITY: CONTAINS 3 CUT DOMAINS.
CC -1- SIMILARITY: BELONGS TO THE CUT FAMILY OF HOMEBOX PROTEINS.
CC -----
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CC -----
DR EMBL; M74099; -; NOT_ANNOTATED_CDS.
DR EMBL; AF047825; AAC78778.1; -;
DR HSSP; P10037; LAU7.
DR TRANSFAC; T00100; -;
DR MIM; 116896; -;
DR InterPro; IPR003350; CUT.
DR InterPro; IPR001356; Homeobox.
DR Pfam; PF02376; CUT; 3.
DR Pfam; PF00046; homeobox; 1.
DR SMART; SM00389; HOX; 1.
DR PROSITE; PS00027; HOMEBOX_1; 1.
DR PROSITE; PSS0071; HOMEBOX_2; 1.
KW Transcription regulation; Homeobox; DNA-binding;
KW Developmental protein; Nuclear protein; Repeat; Repressor;
KW Coiled coil; Alternative splicing.
FT DOMAIN 7 363 COILED COIL (POTENTIAL).
FT DOMAIN 542 629 CUT 1.
FT DOMAIN 669 725 COILED COIL (POTENTIAL).
FT DNA_BIND 934 1021 CUT 2.
FT DNA_BIND 1117 1204 CUT 3.
FT DNA_BIND 1244 1303 HOMEBOX.
FT VARSPIC 632 653 MISSING (IN ISOFORM 2).
SQ SEQUENCE 1505 AA; 164353 MW; 860E14D508D4DE11 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1505;
Best Local Similarity 66.7%; Pred. No. 1.2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
| | | | |
Db 499 EAGSTS 504

RESULT 39
UGG_DROME
ID UGG_DROME STANDARD; PRT; 1548 AA.
AC Q09332;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE UDP-glucose:glycoprotein glucosyltransferase precursor (EC 2.4.1.-)

DE (UDP-Glc:glycoprotein glucosyltransferase) (dugt).
GN UGT OR UGGG.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydriidae; Drosophilidae; Drosophila.
OX NCBI_Taxid=7227;
RN [1]
RP SEQUENCE FROM N.A., AND SEQUENCE OF 23-37.
RC TISSUE=Embryo;
RX MEDLINE=95246722; PubMed=7729408;
RA Parker C.G., Fessler L.I., Nelson R.E., Fessler J.H.;
RT "Drosophila UDP-glucose:glycoprotein glucosyltransferase: sequence
and characterization of an enzyme that distinguishes between
denatured and native proteins.";
RL EMBO J. 14:1294-1303(1995).
CC -!- FUNCTION: UNFOLDED, DENATURED GLYCOPROTEINS ARE SUBSTANTIALLY
CC BETTER SUBSTRATES FOR GLUCOSYLATION BY THIS ENZYME THAN ARE THE
CC CORRESPONDING NATIVE PROTEINS. THIS PROTEIN AND TRANSIENT
CC GLUCOSYLATION MAY BE INVOLVED IN MONITORING AND/OR ASSISTING THE
CC FOLDING AND ASSEMBLY OF NEWLY MADE GLYCOPROTEINS, IN ORDER TO
CC IDENTIFY GLYCOPROTEINS THAT NEED ASSISTANCE IN FOLDING FROM
CC CHAPERONES.
CC -!- COFACTOR: REQUIRES CALCIUM AND MANGANESE IONS FOR ACTIVITY.
CC -!- PATHWAY: GLUCOSYLATION.
CC -!- SUBUNIT: MONOMER.
CC -!- SUBCELLULAR LOCATION: Endoplasmic reticulum.
CC -!- DEVELOPMENTAL STAGE: IS PRESENT AT LOW BUT DETECTABLE LEVELS IN
CC THE EARLIEST EMBRYOS, INCREASING AT 6-8 HRS WITH A MAXIMUM AT 10-
CC 12 HRS. LEVELS DECREASE THEREAFTER AND ARE NOT DETECTED IN 18-20
CC HRS EMBRYOS AND FIRST INSTAR LARVAE BUT IS DETECTED AGAIN AT
CC SECOND INSTAR TO PUPATION.
CC -!- SIMILARITY: SOME, TO YEAST KRE5, S.TYPHIMURIUM RFAJ AND E.COLI
CC RFAI PROTEINS.
CC -----
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CC -----
DR EMBL; U20554; AAA85850.1; -;
DR FlyBase; FBgn0014075; Ugt.
DR InterPro; IPR002495; Glycosyl_transf_8.
DR Pfam; PF01501; Glyco_transf_8; 1.
KW Signal; Transferase; Glycosyltransferase; Endoplasmic reticulum;
KW Glycoprotein. 1 22
FT CHAIN 23 1548 UDP-GLUCOSE:GLYCOPROTEIN
FT GLYCOSYLTRANSFERASE.
FT CARBOHYD 181 181 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 266 266 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 864 864 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT SITE 1545 1548 PREVENT SECRETION FROM ER (POTENTIAL).
SQ SEQUENCE 1548 AA; 174465 MW; 95D6849961622DB6 CRC64;

Query Match 90.5%; Score 19; DB 1; Length 1548;
Best Local Similarity 66.7%; Pred. No. 1.3e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
||| |
Db 255 EAGSTS 260

RESULT 40
HMP2_YEREN STANDARD; PRT; 2035 AA.
AC P48633;
DT 01-FEB-1996 (Rel. 33, Created)

DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE High-molecular-weight protein 2 (HMPW2).
GN IRP2
OS Versinia enterocolitica.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Versinia.
OX NCBI_Taxid=630;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=8081 / SEROTYPE O:8;
RX MEDLINE=93374844; PubMed=8366034;
RA Guilvout I., Mercereau-Puijalon O., Bonnefoy S., Pugsley A.P.,
RA Carniel E.;
RT "High-molecular-weight protein 2 of Versinia enterocolitica is
homologous to AngR of Vibrio anguillarum and belongs to a family of
proteins involved in nonribosomal peptide synthesis.";
RL J. Bacteriol. 175:5488-5504(1993).
CC -!- FUNCTION: UNKNOWN. MAY BE INVOLVED IN THE NONRIBOSOMAL SYNTHESIS
CC OF SMALL PEPTIDES.
CC -!- COFACTOR: CONTAINS 3 COVALENTLY BOUND PHOSPHOPANTHETINES
CC (POTENTIAL).
CC -!- DOMAIN: CONSISTS OF A CENTRAL REGION WITH SIMILARITY TO THE REPEAT
CC DOMAINS OF ACVS AND GRC2, FLANKED BY TWO REPEAT DOMAINS, EACH OF
CC WHICH CONTAINS 5 DIRECT REPEATS.
CC -!- SIMILARITY: BELONGS TO THE ATP-DEPENDENT AMP-BINDING ENZYME
CC FAMILY.
CC -!- SIMILARITY: CONTAINS 3 ACYL CARRIER DOMAINS.
CC -----
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CC -----
DR EMBL; L18881; AAA27636.1; -;
DR EMBL; Z35454; CAA84606.1; -;
DR PIR; A48654; A48654.
DR HSP; P14687; LAMU.
DR InterPro; IPR000873; AMP-bind.
DR InterPro; IPR001242; DUF4.
DR InterPro; IPR003880; Phosphopant_attach.
DR InterPro; IPR000051; SAM_bind.
DR Pfam; PF00501; AMP-binding; 1.
DR Pfam; PF00668; Condensation; 2.
DR Pfam; PF00550; pp-binding; 3.
DR PRINTS; PR00154; AMPBINDING.
DR PROSITE; PS00012; PHOSPHOPANTHETINE; 1.
DR PROSITE; PS00455; AMP_BINDING; 1.
DR PROSITE; PS00075; ACP_DOMAIN; 3.
KW Ligase; Multifunctional enzyme; Phosphopantetheine; Repeat.
FT DOMAIN 3 547 I.
FT REPEAT 114 146 I-DR1.
FT REPEAT 310 321 I-DR2.
FT REPEAT 378 390 I-DR3.
FT REPEAT 454 462 I-DR4.
FT REPEAT 477 491 I-DR5.
FT DOMAIN 1466 1919 II.
FT REPEAT 1495 1527 II-DR1.
FT REPEAT 1682 1693 II-DR2.
FT REPEAT 1750 1762 II-DR3.
FT REPEAT 1826 1834 II-DR4.
FT REPEAT 1849 1863 II-DR5.
FT DOMAIN 20 88 ACYL CARRIER (ACP) 1.
FT DOMAIN 1409 1475 ACYL CARRIER (ACP) 2.
FT DOMAIN 1944 2014 ACYL CARRIER (ACP) 3.
FT BINDING 52 52 PHOSPHOPANTHETINE (BY SIMILARITY).
SQ SEQUENCE 2035 AA; 228826 MW; 1C801377A4375BDC CRC64;

Query Match

90.5%; Score 19; DB 1; Length 2035;

```

Best Local Similarity 66.7%; Pred. No. 1.7e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
Db 1972 EAGTS 1977.

RESULT 41
CLPA_PNPS STANDARD; PRT; 30 AA.
ID CLPA_PNPS STANDARD; PRT; 30 AA.
AC P81671;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DE ATP-dependent clp protease ATP-binding subunit clpA homolog
DE (Fragments)
OS Pinus pinaster (Maritime pine).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Coniferopsida; Coniferales; Pinaceae; Pinus.
OX NCBI_TaxID=71647;
RN [1]
RP SEQUENCE.
RC TISSUE=Needle;
RX MEDLINE=99274088; PubMed=10344291;
RA Costa P., Pionneau C., Bauw G., Dubos C., Bahrman N., Kremer A.,
RA Frigerio J.-M., Plomion C.;
RT "Separation and characterization of needle and xylem maritime pine
RT proteins";
RL Electrophoresis 20:1098-1108(1999).
CC -1- FUNCTION: MAY INTERACT WITH A CLPP-LIKE PROTEASE INVOLVED IN
CC DEGRADATION OF DENATURED PROTEINS IN THE CHLOROPLAST (BY
CC SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Chloroplast (By similarity).
CC -1- MISCELLANEOUS: ON THE 2D-GEL THE DETERMINED PI OF THIS PROTEIN
CC (SPOT N9) IS: 5.9, ITS MW IS: 92 kDa.
CC -1- SIMILARITY: BELONGS TO THE CLPA/CLPB FAMILY.
DR InterPro; IPR001270; CLP_AB.
DR PROSITE; PS00870; CLPAB_1; PARTIAL.
DR PROSITE; PS00871; CLPAB_2; PARTIAL.
KW Chapterone; ATP-binding; Repeat; Chloroplast.
FT NON_TER 1 1
FT NON_CONS 15 16
FT NON_TER 30 30
FT NON_CONS 30 30
SQ SEQUENCE 30 AA; 2923 MW; 44B5950B73A96152 CRC64;

Query Match 85.7%; Score 18; DB 1; Length 30;
Best Local Similarity 66.7%; Pred. No. 47;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
Db 4 EAGDAS 9

RESULT 42
PLAS_CAPBU STANDARD; PRT; 99 AA.
ID PLAS_CAPBU STANDARD; PRT; 99 AA.
AC P00294;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Plastocyanin.
DE Plastocyanin.
GN PTE.
OS Capsella bursa-pastoris (Shepherd's purse).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Capsella.
OX NCBI_TaxID=3719;
RN [1]
RP SEQUENCE.
RA Scawen M.D., Ramshaw J.A.M., Brown R.H., Boulter D.;

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Unpublished results, cited by:
Boulter D., Haslett B.G., Peacock D., Ramshaw J.A.M., Scawen M.D.;
(In) Northcote D.H. (eds.);
Plant biochemistry II, pp.13:1-40, University Park Press,
Baltimore (1977).
CC -1- FUNCTION: PLASTOCYANIN PARTICIPATES IN ELECTRON TRANSFER BETWEEN
CC P700 AND THE CYTOCHROME B/F COMPLEX IN PHOTOSYSTEM I.
CC -1- SUBCELLULAR LOCATION: LOOSELY BOUND TO THE INNER THYLAKOID
CC MEMBRANE SURFACE IN CHLOROPLASTS.
CC -1- SIMILARITY: CONTAINS 1 PLASTOCYANIN-LIKE DOMAIN.
DR PIR; A00304; CUSU.
DR HSP; P00289; 2PCF.
DR Mendel; I1575; CAPBU; PteI.1.
DR InterPro; IPR001235; Copper_blue.
DR InterPro; IPR000923; Copper_blue.
DR Pfam; PF00127; copper-bind; 1.
DR PRINTS; PR00156; COPPERBLUE.
DR ProDom; PD001235; Copper_blue; 1.
DR PROSITE; PS00196; COPPER_BLUE; 1.
KW Chloroplast; Electron transport; Copper; Thylakoid; Membrane.
FT DOMAIN 1 99 PLASTOCYANIN-LIKE.
FT METAL 37 37 COPPER (BY SIMILARITY).
FT METAL 84 84 COPPER (BY SIMILARITY).
FT METAL 87 87 COPPER (BY SIMILARITY).
FT METAL 92 92 COPPER (BY SIMILARITY).
SQ SEQUENCE 99 AA; 10383 MW; 30BA97B58B9580F1 CRC64;

Query Match 85.7%; Score 18; DB 1; Length 99;
Best Local Similarity 66.7%; Pred. No. 1.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
Db 76 EAGYS 81

RESULT 43
WN14_HUMAN STANDARD; PRT; 123 AA.
ID WN14_HUMAN STANDARD; PRT; 123 AA.
AC O14904;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE WNT-14 protein (fragment).
GN WNT14
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98110581; PubMed=9441749;
RA Bergstein I., Eisenberg L.M., Bhalarao J., Jenkins N.A.;
RA Copeland N.G., Osborne M.P., Bowcock A.M., Brown A.M.C.;
RT "Isolation of two novel WNT genes, WNT14 and WNT15, one of which
RT (WNT15) is closely linked to WNT3 on human chromosome 17q21.";
RL Genomics 46:450-458(1997).
CC -1- FUNCTION: LIGAND FOR MEMBERS OF THE FRIZZLED FAMILY OF SEVEN
CC TRANSMEMBRANE RECEPTORS. PROBABLE DEVELOPMENTAL PROTEIN. MAY BE A
CC SIGNALING MOLECULE WHICH AFFECT THE DEVELOPMENT OF DISCRETE
CC REGIONS OF TISSUES. IS LIKELY TO SIGNAL OVER ONLY FEW CELL
CC DIAMETERS (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Possibly secreted and associates with the
CC extracellular matrix.
CC -1- SIMILARITY: BELONGS TO THE WNT FAMILY.
CC -----
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CC -----
DR EMBL; AF028702; AAC39550.1; -.
DR MIM; 602863; -.
DR InterPro; IPR000970; Wnt1.
DR Pfam; PF00110; wnt; 1.
DR SMART; SM00097; WNT1; 1.
DR PROSITE; PS00246; WNT1; PARTIAL.
KW Developmental protein; Glycoprotein.
FT NON_TER 1
FT NON_TER 123
SQ SEQUENCE 123 AA; 13143 MW; 8F000D2568EEA744 CRC64;

Query Match
Best Local Similarity 85.7%; Score 18; DB 1; Length 123;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
DB 43 EAGAIS 48

RESULT 44
PPIB_BACSU
ID PPIB_BACSU STANDARD; PRT; 143 AA.
AC P35137;
DT 01-FEB-1994 (Rel. 28, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Peptidyl-prolyl cis-trans isomerase B (EC 5.2.1.8) (PPIase B)
DE (Rotamase B).
GN PPIB.
OS Bacillus subtilis.
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Bacillus.
OX NCBI_TaxID=1423;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=168 / MAREBURG;
RX MEDLINE=95020538; PubMed=7934829;
RA Sorokin A.V., Zumbstein E., Azevedo V., Ehrlich S.D., Serrero P.;
RT "The organization of the Bacillus subtilis 168 chromosome region
between the spoVA and serA genetic loci, based on sequence data.";
RL Mol. Microbiol. 10:385-395(1993).
RN [2]
RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
RC STRAIN=168 / JH642;
RX MEDLINE=94293776; PubMed=8022278;
RA Herrler M., Bang H., Marahiel M.A.;
RT "Cloning and characterization of ppiB, a Bacillus subtilis gene which
encodes a cyclosporin A-sensitive peptidyl-prolyl cis-trans
isomerase.";
RT isomerase.";
RL Mol. Microbiol. 11:1073-1083(1994).
RN [3]
RP SEQUENCE OF 1-26.
RC STRAIN=168 / JH642;
RX MEDLINE=96345629; PubMed=8755892;
RA Graumann P., Schroeder K., Schmid R., Marahiel M.A.;
RT "Cold shock stress-induced proteins in Bacillus subtilis.";
RL J. Bacteriol. 178:4611-4619(1996).
CC -1- FUNCTION: PPIASES ACCELERATE THE FOLDING OF PROTEINS.
CC -1- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC
PEPTIDE BONDS IN OLIGOPEPTIDES.
CC -1- ENZYME REGULATION: INHIBITED BY CYCLOSPORIN A (CSA).
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -1- SIMILARITY: BELONGS TO THE CYCLOPHILIN-TYPE PPIASE FAMILY.
CC -----
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CC -----
CC EMBL; L09228; AAA67475.1; -.
CC EMBL; X73898; CAA52103.1; -.
CC EMBL; Z99116; CABI4288.1; -.
CC PIR; S45537; S45537.
CC HSP; Q27450; 1A58.
CC Subtilisin; BG10512; ppiB.
CC InterPro; IPR002130; CSA_PPIase.
CC Pfam; PF00160; pro_isomerase; 1.
CC PRINTS; PR00153; CSAPPISMRASE.
CC PROSITE; PS00170; CSA_PPIASE_1; 1.
CC PROSITE; PS00072; CSA_PPIASE_2; 1.
KW Isomerase; Rotamase; Complete proteome.
SQ SEQUENCE 143 AA; 15256 MW; 9EF17D70EB81EC51 CRC64;

Query Match
Best Local Similarity 85.7%; Score 18; DB 1; Length 143;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
DB 82 EAGALS 87

RESULT 45
R157_BOVIN
ID R157_BOVIN STANDARD; PRT; 147 AA.
AC Q28183;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE Retina-specific 15.7 kDa protein.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Retina;
RX MEDLINE=86258168; PubMed=2425311;
RA Nakagawa Y., Kuo C.H., Ishii K., Shiosaka S., Tohyama M., Miki N.;
RT "Cloning and characterization of a cDNA specific for bovine retina.";
RL Neurosci. Res. 3:300-310(1986).
CC -1- TISSUE SPECIFICITY: RETINA.
CC -----
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CC -----
CC EMBL; M34915; AAA30756.1; -.
CC SEQUENCE 147 AA; 15658 MW; 1FEDA4878B39645 CRC64;

Query Match
Best Local Similarity 85.7%; Score 18; DB 1; Length 147;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
DB 133 EAGTVS 138

RESULT 46
DUT_MYCTU
ID DUT_MYCTU STANDARD; PRT; 154 AA.
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AC 007199;
DT 30-MAY-2000 (Rel. 39, Last created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Deoxyuridine 5'-triphosphate nucleotidohydrolase (EC 3.6.1.23)
DE (dUTPase) (dUTP pyrophosphatase).
GN DUT OR RV2697C OR MT2771 OR MTCV05A6.18C.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RX MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekaiia F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544 (1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J.F., DeBoy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
RA Bishai W.;
RA "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: THIS ENZYME IS INVOLVED IN NUCLEOTIDE METABOLISM: IT
CC PRODUCES DUMP, THE IMMEDIATE PRECURSOR OF THYMIDINE NUCLEOTIDES
CC AND IT DECREASES THE INTRACELLULAR CONCENTRATION OF DUTP SO THAT
CC URACIL CANNOT BE INCORPORATED INTO DNA (BY SIMILARITY).
CC -1- CATALYTIC ACTIVITY: dUTP + H(2)O -> dUMP + diphosphate.
CC -1- PATHWAY: DE NOVO SYNTHESIS OF THYMIDYLATE.
CC -1- SIMILARITY: BELONGS TO THE DUTPASE FAMILY.
CC -----
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Query Match 85.7%; Score 18; DB 1; Length 154;
Best Local Similarity 66.7%; Pred. No. 2.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Qy 1 eagxxs 6
Db 132 EAGLAS 137
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RESULT 47
MOAE_RHIME
ID MOAE_RHIME STANDARD; PRT; 155 AA.
AC Q92QX5; 2002 (Rel. 41, Created)
DT 01-MAR-2002 (Rel. 41, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Molybdopterin synthase cofactor subunit 2 (MPT synthase subunit 2)
DE (Molybdopterin synthase subunit 2) (Molybdenum cofactor biosynthesis
DE protein E) (Molybdopterin converting factor large subunit).
GN MOAE OR R01168 OR SMC00599.
OS Rhizobium meliloti (Sinorhizobium meliloti).
OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
OC Rhizobiaceae; Sinorhizobium.
OX NCBI_TaxID=382;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=1021;
RX MEDLINE=21396507; PubMed=11481430;
RA Capela D., Barloy-Hubler F., Gouzy J., Bothe G., Ampe F., Batut J.,
RA Boistard P., Becker A., Boutry M., Cadieu E., Dreano S., Gloux S.,
RA Godrie T., Goffeau A., Kahn D., Kiss E., Lelaure V., Masuy D.,
RA Pohl T., Portetelle D., Puehler A., Purnelle B., Ramsperger G.,
RA Renard C., Thebaud P., Vandenbol M., Weidner S., Galibert F.;
RA "Analysis of the chromosome sequence of the legume symbiont
RT Sinorhizobium meliloti strain 1021.";
RN Proc. Natl. Acad. Sci. U.S.A. 98:9877-9882 (2001).
RL CC -1- FUNCTION: Converts molybdopterin precursor Z into molybdopterin.
CC This requires the incorporation of two sulfur atoms into precursor
CC Z to generate a dithiolene group (By similarity).
CC -1- PATHWAY: Molybdenum cofactor biosynthesis.
CC -1- SUBUNIT: Heterodimer of 2 moad subunits and 2 moae subunits (By
CC similarity).
CC -1- SIMILARITY: BELONGS TO THE MOAE FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
```

RNA SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / HAL;
RX MEDLINE=96079490; PubMed=8566710;
RA Zhao S., Mitchell S.E., Meng J., Doyle M.P., Kresovich S.;
RT "Cloning and nucleotide sequence of a gene upstream of the eaeA gene
of enterohemorrhagic Escherichia coli O157:H7";
RL FEMS Microbiol. Lett. 133:35-39(1995).
[2]
RNA SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / EDL933 / ATCC 700927;
RX MEDLINE=98339885; PubMed=9673266;
RA Perna N.T., Mayhew G.F., Posfal G., Elliott S., Donnenberg M.S.,
R Kaper J.B., Blattner F.R.;
RT "Molecular evolution of a pathogenicity island from enterohemorrhagic
Escherichia coli O157:H7";
RL Infect. Immun. 66:3810-3817(1998).
[3]
RNA SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / ATCC 700927;
RX MEDLINE=21074935; PubMed=11206551;
RA Perna N.T., Plunkett G. III, Burland V., Mau B., Glasner J.D.,
RA Rose D.J., Mayhew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,
RA Postai G., Hackett J., Klink S., Boutin A., Shao Y., Miller L.,
RA Grobeck E.J., Davis N.W., Lim A., Dimalanta E.F., Potamoukis K.,
RA Apodaca J., Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,
RA Welch R.A., Blattner F.R.;
RT "Genome sequence of enterohemorrhagic Escherichia coli O157:H7";
RL Nature 409:529-533(2001).
[4]
RNA SEQUENCE FROM N.A.
RC STRAIN=O157:H7 / RMD 0509952;
RX MEDLINE=21156231; PubMed=11258796;
RA Hayashi T., Makino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,
RA Han C.-G., Ohtsubo E., Nakayama K., Murata T., Tanaka M., Tohe T.,
RA Iida T., Takani H., Honda T., Sasaki C., Ogasawara N., Yasunaga T.,
RA Kuhara S., Shiba T., Hattori M., Shinagawa H.;
RT "Complete genome sequence of enterohemorrhagic Escherichia coli
O157:H7 and genomic comparison with a laboratory strain K-12";
RL DNA Res. 8:11-22(2001).
CC -1- FUNCTION: Chaperone for the type III secretion of Tir. Probably
stabilizes the protein, prevents inappropriate protein-
protein interactions and aids in secretion (By similarity).
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -1- SIMILARITY: BELONGS TO THE CEST/SYCH CHAPERONE FAMILY.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation -
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or send an email to license@isb-sib.ch).

DR EMBL; U32312; AAB00110.1; -;
DR EMBL; AF071034; AAC31505.1; -;
DR EMBL; AE005595; AAG58824.1; -;
DR EMBL; AP002566; BAB37983.1; -;
KW Chaperone; Virulence.
SQ SEQUENCE 156 AA; 17681 MW; 999545426E26D2D6 CRC64;

Query Match 85.7%; Score 18; DB 1; Length 156;
Best Local Similarity 66.7%; Pred. No. 2.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
DB 87 EAGAQS 92

RESULT 49
FLIN_PSEAE

ID FLIN_PSEAE STANDARD; PRT; 157 AA.
AC Q51466;
DT 01-NOV-1997 (Rel. 35, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Flagellar motor switch protein flin.
GN FLIN OR PA1444.
OS Pseudomonas aeruginosa.
OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;
OC Pseudomonas
OX NCBI_TaxID=287;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PAK;
RX MEDLINE=95347807; PubMed=7622217;
RA Simpson D.A., Ramphal R., Lory S.;
RT "Characterization of Pseudomonas aeruginosa fllo, a gene involved in
flagellar biosynthesis and adherence";
RL Infect. Immun. 63:2950-2957(1995).
[2]
RNA SEQUENCE FROM N.A.
RC STRAIN=ATCC 15692 / PA01;
RX MEDLINE=20437337; PubMed=10984043;
RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
RA Hickey M.J., Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,
RA Garber R.L., Goltry L., Tolentino E., Westbrook-Wadman S., Yuan Y.,
RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
RA Reizer J., Saler M.H., Hancock R.E.W., Lory S., Olson M.V.;
RT "Complete genome sequence of Pseudomonas aeruginosa PA01, an
opportunistic pathogen";
RL Nature 406:959-964(2000).
CC -1- FUNCTION: FLIN IS ONE OF THREE PROTEINS (FLIG, FLIN, FLIM) THAT
FORM A SWITCH COMPLEX THAT IS PROPOSED TO BE LOCATED AT THE BASE
OF THE BASAL BODY. THIS COMPLEX INTERACTS WITH THE CHEY AND CHEZ
CHEMOTAXIS PROTEINS, IN ADDITION TO CONTACTING COMPONENTS OF THE
MOTOR THAT DETERMINE THE DIRECTION OF FLAGELLAR ROTATION (BY
SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Inner membrane-associated (Potential).
CC -1- SIMILARITY: BELONGS TO THE FLIN/MOPA/SPAO FAMILY.
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entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
or send an email to license@isb-sib.ch).

DR EMBL; L39832; AAA79754.1; -;
DR EMBL; AE004574; AAG04833.1; -;
DR InterPro: IPR001172; Flag_Flin.
DR InterPro: IPR001543; Spoa.
DR Pfam: PF01052; Spoa; 1.
DR PRINTS; PR00956; FLGMOTORFLIN.
DR ProDom; PD001777; Spoa; 1.
KW Chemotaxis; Flagella; Flagellar rotation; Inner membrane;
KW Complete proteome.
FT CONFLICT 48 48 P -> S (IN REF. 1).
SQ SEQUENCE 157 AA; 16620 MW; B3D91C0182ACB775 CRC64;

Query Match 85.7%; Score 18; DB 1; Length 157;
Best Local Similarity 66.7%; Pred. No. 2.5e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
DB 24 EAGDAS 29

RESULT 50
YMH2_CAEEL

```
ID YMH2_CAEEL STANDARD; PRT; 159 AA.
AC P34469;
DT 01-FEB-1994 (Rel. 28, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Hypothetical 17.2 kDa protein F58A4.2 in chromosome III.
GN F58A4.2.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RX MEDLINE=94150718; PubMed=7906398;
RA Wilson R., Alnsough R., Anderson K., Baynes C., Berks M.,
RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
RA Craxton M., Dear S., Du Z., Durbin R., Favello A., Fraser A.,
RA Fulton L., Gardner A., Green P., Hawkins T., Hillier L., Jier M.,
RA Johnston L., Jones M., Kershaw J., Kirsten J., Laisster N.,
RA Latreille P., Lightning J., Lloyd C., Mortimore B., O'Callaghan M.,
RA Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Showkeen R.,
RA Sims M., Smaldon N., Smith A., Smith M., Sonnhammer E., Staden R.,
RA Sulston J., Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K.,
RA Waterston R., Watson A., Weinstock L., Wilkinson-Sproat J.,
RA Wohlman P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans.";
RL Nature 368:32-38(1994).
CC -----
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC -----
DR EMBL; Z22179; CAA80168.1; -.
DR PIR; S40974; S40974.
DR WormPep; F58A4.2; CE01017.
KW Hypothetical protein.
SQ SEQUENCE 159 AA; 17201 MW; 364FE35A65E2C89D CRC64;
```

```
Query Match 85.7%; Score 18; DB 1; Length 159;
Best Local Similarity 66.7%; Pred. No. 2.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
```

```
Qy 1 eagxxs 6
   ||| |
Db 145 EAGSGS 150
```

Search completed: August 30, 2002, 15:11:55
Job time: 311 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: August 30, 2002, 15:05:49 ; Search time 41.29 Seconds
(without alignments)
25.139 Million cell updates/sec

Title: BASK-853-CLAIM4

Perfect score: 21
Sequence: 1 eagxxs 6

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 562222 seqs, 172994929 residues

Total number of hits satisfying chosen parameters: 562222

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 50 summaries

Database : SPTREMBL_19.*
1: sp_archaea.*
2: sp_bacteria.*
3: sp_fungi.*
4: sp_human.*
5: sp_invertebrate.*
6: sp_mammal.*
7: sp_mhc.*
8: sp_organelle.*
9: sp_phage.*
10: sp_plant.*
11: sp_rodent.*
12: sp_virus.*
13: sp_vertebrate.*
14: sp_unclassified.*
15: sp_virus.*
16: sp_bacteriap.*
17: sp_archaeap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19	90.5	108	12 Q91TM3	Q91tm3 tupaia herp
2	19	90.5	122	17 Q9HPE3	Q9hpe3 halobacteri
3	19	90.5	130	15 Q36890	Q36890 human immun
4	19	90.5	131	6 Q9GK47	Q9gk47 galago cras
5	19	90.5	143	17 Q9VXB6	Q9vxb6 aeropyrum p
6	19	90.5	145	16 Q9KR08	Q9kr08 vibrio chol
7	19	90.5	149	10 Q9FRJ5	Q9frj5 oryza sativ
8	19	90.5	151	10 Q9AW19	Q9aw19 oryza sativ
9	19	90.5	152	10 Q9A1K5	Q9a1k5 arabidopsis
10	19	90.5	161	4 Q9BS09	Q9bs09 homo sapien
11	19	90.5	161	4 Q9NWI1	Q9nwi1 homo sapien
12	19	90.5	164	2 Q54209	Q54209 streptomyce
13	19	90.5	170	6 Q9WZ11	Q9wz11 oryctolagus
14	19	90.5	192	10 Q94L73	Q94l73 oryza sativ
15	19	90.5	200	5 Q18144	Q18144 caenorhabdi
16	19	90.5	201	5 Q62323	Q62323 caenorhabdi

17	19	90.5	204	16 Q53374	Q53374 mycobacteri
18	19	90.5	212	16 Q92D00	Q92d00 listeria in
19	19	90.5	231	13 Q91306	Q91306 rana catesb
20	19	90.5	245	17 Q9HST1	Q9hst1 halobacteri
21	19	90.5	254	13 Q91307	Q91307 rana catesb
22	19	90.5	257	10 Q9XF64	Q9xf64 arabidopsis
23	19	90.5	257	10 Q9LZJ6	Q9lzf6 arabidopsis
24	19	90.5	260	16 Q99RZ3	Q99rz3 staphylococ
25	19	90.5	261	2 Q9ZIN7	Q9zin7 staphylococ
26	19	90.5	261	16 Q92NH7	Q92nh7 rhizobium m
27	19	90.5	266	12 Q88190	Q88190 soybean mos
28	19	90.5	267	12 Q88196	Q88196 soybean mos
29	19	90.5	269	11 Q9DCE4	Q9dce4 mus musculu
30	19	90.5	270	5 Q9VGK4	Q9vgk4 drosophila
31	19	90.5	285	2 Q9A1T5	Q9ait5 vibrio chol
32	19	90.5	287	13 Q93503	Q93503 xenopus lae
33	19	90.5	293	2 Q91866	Q91866 azospirillu
34	19	90.5	297	13 Q91296	Q91296 rana catesb
35	19	90.5	302	16 Q9PCJ6	Q9pcj6 xyliella fas
36	19	90.5	307	10 Q9FJG2	Q9fjg2 arabidopsis
37	19	90.5	311	13 Q90888	Q90888 gallus gall
38	19	90.5	311	13 Q90370	Q90370 coturnix co
39	19	90.5	313	13 Q9PUA6	Q9puae xenopus lae
40	19	90.5	323	4 Q9Y5Q3	Q9y5q3 homo sapien
41	19	90.5	323	4 Q9H1F1	Q9h1f1 homo sapien
42	19	90.5	333	13 Q98TS3	Q98ts3 brachydanio
43	19	90.5	346	10 Q9ATS6	Q9ats6 arundinella
44	19	90.5	350	16 Q981H9	Q981h9 rhizobium l
45	19	90.5	352	16 Q92WT2	Q92wt2 rhizobium m
46	19	90.5	356	13 Q98UK5	Q98uk5 brachydanio
47	19	90.5	356	13 Q73679	Q73679 brachydanio
48	19	90.5	357	10 Q23101	Q23101 arabidopsis
49	19	90.5	361	5 Q9BL91	Q9bl91 caenorhabdi
50	19	90.5	362	11 Q9JHQ1	Q9jhq1 rattus norv

ALIGNMENTS

RESULT 1
Q91TM3 PRELIMINARY; PRT; 108 AA.
AC Q91TM3;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE 773.
OS Tupaia herpesvirus.
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Betaherpesvirinae.
OX NCBI_TaxID=10397;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=2;
RX MEDLINE=2111637; PubMed=11312357;
RA Bahr U., Darai G.;
RT "Analysis and Characterization of the Complete Genome of Tupaia (Tree Shrew) Herpesvirus".
RL J. Virol. 75:4854-4870(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=2;
RA Darai G., Bahr U.;
RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF281817; AAK57118.1; -.
SQ SEQUENCE 108 AA; 1118 MW; 5732B2C61DBDE820 CRC64;

Query Match 90.5%; Score 19; DB 12; Length 108;

Best Local Similarity 66.7%; Pred. No. 5.8e+02; Indels 0; Gaps 0;
Matches 4; Conservative 0; Mismatches 2;

QY 1 eagxxs 6

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DB 30 EAGASS 35

RESULT 2
Q9HPE3 PRELIMINARY; PRT; 122 AA.
AC Q9HPE3;
DT 01-MAR-2001 (TRENBLrel. 16, Created)
DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
DT 01-MAR-2001 (TRENBLrel. 16, Last annotation update)
DE VNC1678H.
GN VNC1678H.
OS Halobacterium sp. (strain NRC-1).
OC Archaea; Euryarchaeota; Halobacteriales; Halobacteriaceae;
OC Halobacterium.
OX NCBI_TaxID=64091;
RN [1]
RX SEQUENCE FROM N.A.
RX MEDLINE=20504483; PubMed=11016950;
RA Ng W.V., Kennedy S.P., Mahairas G.G., Berquist B., Pan M.,
RA Shukla H.D., Lasky S.R., Balliga N.S., Thorsson V., Sbrogna J.,
RA Swartzell S., Weir D., Hall J., Dahl T.A., Welti R., Goo Y.A.,
RA Leithausen B., Keller K., Cruz R., Danson M.J., Hough D.W.,
RA Maddocks D.G., Jablonski P.E., Krebs M.P., Angevine C.M., Dale H.,
RA Isenbarger T.A., Peck R.F., Pohlschroder M., Spudich J.L., Jung K.-H.,
RA Alam M., Freitas T., Hou S., Daniels C.J., Dennis P.P., Omer A.D.,
RA Ehardt H., Lowe T.M., Liang P., Riley M., Hood L., Dassarma S.;
RT "Genome sequence of Halobacterium species NRC-1.";
RL Proc. Natl. Acad. Sci. U.S.A. 97:12176-12181(2000).
DR EMBL; AE005076; AAG19927.1; -.
KW Complete proteome.
SQ SEQUENCE 122 AA; 12001 MW; BE2432416C2BA256 CRC64;

Query Match 90.5%; Score 19; DB 17; Length 122;
Best Local Similarity 66.7%; Pred. No. 6.6e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
   ||| |
   57 EAGASS 62

DB 57 EAGASS 62

RESULT 3
O36890 PRELIMINARY; PRT; 130 AA.
AC O36890;
DT 01-JAN-1998 (TRENBLrel. 05, Created)
DT 01-JAN-1998 (TRENBLrel. 05, Last sequence update)
DT 01-DEC-2001 (TRENBLrel. 19, Last annotation update)
DE GAG POLYPROTEIN (FRAGMENT).
GN GAG.
OS Human immunodeficiency virus type 1.
OC Viruses; Retroid viruses; Retroviridae; Lentivirus.
OX NCBI_TaxID=11676;
RN [1]
RX SEQUENCE FROM N.A.
RX MEDLINE=97445059; PubMed=9300048;
RA Leigh Brown A.J., Lobidel D., Wade C.M., Rebus S., Phillips N.,
RA Brettie R.P., France A.J., Leen C.S., McMenamin J., McMillan A.,
RA Maw R.D., Mulcahy F., Robertson J.R., Sankar K.N., Scott G., Wyld R.,
RA Peutherer J.F.;
RT "The molecular epidemiology of human immunodeficiency virus type 1 in
RT six cities in Britain and Ireland.";
RL Virology 235:166-177(1997).
DR EMBL; AF014287; AAC58368.1; -.
DR Incerpro; IFP000071; Retroviral_gag_p17.
DR Pfam; PF00540; gag_p17; 1.
DR PRINTS; PR00234; HIVMATRIX.
KW AIDS; Core protein; Polyprotein.
FT NON_TER 1 130
```

```
SQ SEQUENCE 130 AA; 14476 MW; 9053DD2EF3A3E00F CRC64;

Query Match 90.5%; Score 19; DB 15; Length 130;
Best Local Similarity 66.7%; Pred. No. 7.1e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
   ||| |
   99 EAGSSS 104

DB 99 EAGSSS 104

RESULT 4
Q9GK47 PRELIMINARY; PRT; 131 AA.
AC Q9GK47;
DT 01-MAR-2001 (TRENBLrel. 16, Created)
DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
DT 01-DEC-2001 (TRENBLrel. 19, Last annotation update)
DE RELAXIN-LIKE PROTEIN.
OS Galago crassicaudatus (Thick-tailed galago) (Otolemur crassicaudatus).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Strepsirhini; Galagonidae; Otolemur.
OX NCBI_TaxID=9463;
RN [1]
RX SEQUENCE FROM N.A.
RX TISSUE=TESTIS;
RA Klonisch T., Froehlich C., Tetens F., Fischer B., Hombach-Klonisch S.;
RT "Molecular remodeling of members of the relaxin family during primate
RT evolution.";
RL Mol. Biol. Evol. 0:0-0(2001).
CC -!- SUBCELLULAR LOCATION: SECRETED (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE INSULIN/IGF/RELAXIN FAMILY.
DR EMBL; AF317624; AAG42317.1; -.
DR InterPro; IPR000739; Insulin_IGF_relaxin.
DR SMART; SM00078; ILGF; 1.
DR PROSITE; PS00262; INSULIN; 1.
SQ SEQUENCE 131 AA; 14414 MW; F18AFA9ACFC85943 CRC64;

Query Match 90.5%; Score 19; DB 6; Length 131;
Best Local Similarity 66.7%; Pred. No. 7.1e+02;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
   ||| |
   54 EAGTSS 59

DB 54 EAGTSS 59

RESULT 5
Q9YBX6 PRELIMINARY; PRT; 143 AA.
AC Q9YBX6;
DT 01-NOV-1999 (TRENBLrel. 12, Created)
DT 01-NOV-1999 (TRENBLrel. 12, Last sequence update)
DT 01-MAR-2001 (TRENBLrel. 16, Last annotation update)
DE HYPOTHETICAL 15.7 KDA PROTEIN APE1474.
GN APE1474.
OS Aeropyrum pernix.
OC Archaea; Crenarchaeota; Desulfurococcales; Desulfurococcaceae;
OC Aeropyrum.
OX NCBI_TaxID=56636;
RN [1]
RX SEQUENCE FROM N.A.
RX STRAIN=K1;
RA Kwarabayasi Y., Hino Y., Horikawa H., Yamazaki S., Haikawa Y.,
RA Jin-no K., Takahashi M., Sekine M., Baba S.-I., Ankal A., Kosugi H.,
RA Hosoyama A., Fukui S., Nagai Y., Nishijima K., Nakazawa H.,
RA Takamiya M., Masuda S., Funahashi T., Tanaka T., Kudoh Y.,
RA Yamazaki J., Kishida N., Oguchi A., Aoki K.-I., Kubota K.,
RA Nakamura Y., Nomura N., Sako Y., Kikuchi H.;
RT "Complete genome sequence of an aerobic hyper-thermophilic
```

RT crenarchaeon, Aeropyrum pernix K1.";
 RL DNA Res. 6:83-101(1999).
 DR EMBL; AF000061; BAA80472.1; -.
 KW Hypothetical protein; Complete proteome.
 SQ SEQUENCE 143 AA; 15708 MW; C40E29CBB0AF6892 CRC64;

Query Match 90.5%; Score 19; DB 17; Length 143;
 Best Local Similarity 66.7%; Pred. No. 7.8e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
 |||||
 Db 85 EAGAAS 90

RESULT 6
 Q9KR08 PRELIMINARY; PRT; 145 AA.
 AC Q9KR08;
 DT 01-OCT-2000 (TREMBLrel. 15, Created)
 DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
 DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
 DE HYPOTHETICAL PROTEIN VC1536.
 GN VC1536.
 OS Vibrio cholerae.
 OC Bacteria; Proteobacteria; gamma subdivision; Vibrionaceae; Vibrio.
 OX NCBI_TaxID=666;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=EL TOR N16961 / SEROTYPE O1;
 RX MEDLINE=20406833; PubMed=10952301;
 RA Heidelberg J.F., Eisen J.A., Nelson W.C., Clayton R.A., Gwinn M.L.,
 RA Dodson R.J., Haft D.H., Hickey E.K., Peterson J.D., Umayam L.A.,
 RA Gill S.R., Nelson K.E., Read T.D., Tettelin H., Richardson D.,
 RA Ermolaeva M.D., Vamathevan J., Bass S., Qin H., Dragol I., Sellers P.,
 RA McDonald L., Utterback T., Fleischmann R.D., Nierman W.C., White O.,
 RA Salzberg S.L., Smith H.O., Colwell R.R., Mekalanos J.J., Venter J.C.,
 RA Fraser C.M.;
 RT "DNA sequence of both chromosomes of the cholera pathogen Vibrio
 cholerae.";
 RL Nature 406:477-483(2000).
 DR EMBL; AE004231; AAF94690.1; -.
 DR TIGR; VC1536; -.
 KW Hypothetical protein; Complete proteome.
 SQ SEQUENCE 145 AA; 15726 MW; 976E1F5EB50DB0EC CRC64;

Query Match 90.5%; Score 19; DB 16; Length 145;
 Best Local Similarity 66.7%; Pred. No. 7.9e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
 |||||
 Db 114 EAGSTS 119

RESULT 7
 Q9FRJ5 PRELIMINARY; PRT; 149 AA.
 AC Q9FRJ5;
 DT 01-MAR-2001 (TREMBLrel. 16, Created)
 DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
 DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
 DE HYPOTHETICAL 15.3 KDA PROTEIN.
 GN OSJNB0064P21.9.
 OS Oryza sativa (rice).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Ehrhartoideae; Oryzeae; Oryza.
 OX NCBI_TaxID=4530;
 RN [1]
 RP SEQUENCE FROM N.A.

RC STRAIN=CV. NIPPONBARE;
 RA Buell C.R., Yuan Q., Moffat K.S., Hill J.N., Burr P.C., Hsiao J.,
 RA Zismann V., Pai G., Bowman C.L., Fujii C.Y., VanAken S.E.,
 RA Bowman C.L., Craven B., Utterback T.R., Khalak H., Feldblyum T.V.,
 RA Quackenbush J., White O., Salzberg S.L., Fraser C.M.;
 RT "Oryza sativa chromosome 10 BAC OSJNB0064P21 genomic sequence.";
 RL Submitted (AUG-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AC073166; AAG46108.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 149 AA; 15284 MW; 8B71E92310872766 CRC64;

Query Match 90.5%; Score 19; DB 10; Length 149;
 Best Local Similarity 66.7%; Pred. No. 8.1e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
 |||||
 Db 111 EAGASS 116

RESULT 8
 Q9AWI9 PRELIMINARY; PRT; 151 AA.
 ID Q9AWI9;
 AC Q9AWI9;
 DT 01-JUN-2001 (TREMBLrel. 17, Created)
 DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE P0489A05.8 PROTEIN.
 GN P0489A05.8.
 OS Oryza sativa (rice).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Ehrhartoideae; Oryzeae; Oryza.
 OX NCBI_TaxID=4530;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. NIPPONBARE;
 RA Sasaki T., Matsumoto T., Yamamoto K.;
 RT "Oryza sativa nipponbare(GA3) genomic DNA, chromosome 1, PAC
 clone:P0489A05.";
 RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AF003105; BAB32988.1; -.
 SQ SEQUENCE 151 AA; 16103 MW; 6E942A203BC62C11 CRC64;

Query Match 90.5%; Score 19; DB 10; Length 151;
 Best Local Similarity 66.7%; Pred. No. 8.2e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eaqxss 6
 |||||
 Db 106 EAGAAS 111

RESULT 9
 Q9XIK5 PRELIMINARY; PRT; 152 AA.
 ID Q9XIK5;
 AC Q9XIK5;
 DT 01-NOV-1999 (TREMBLrel. 12, Created)
 DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)
 DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
 DE T10024.6.
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
 OC eurosids II; Brassicales; Brassicaceae; Arabidopsi.
 OX NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.

RA Shinn P., Brooks S., Buehler E., Chao Q., Dunn P., Khan S., Kim C.,
 RA Walker M., Altafi H., Araujo R., Conn L., Conway A.B., Gonzalez A.,
 RA Hansen N.F., Huizar L., Kremenetska I., Lenz C., Li J., Liu S.,

RA Lueros S., Rowley D., Schwartz J., Toriumi M., Vysotskaia V., Yu G.,
 RA Davis R.W., Federspiel N.A., Theologis A., Ecker J.R.;
 RT "Genomic sequence for Arabidopsis thaliana BAC T10024 from Chromosome
 1.";
 RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AC007067; AAD39566.1; -.
 SQ SEQUENCE 152 AA; 17676 MW; A7053F4DA73C3490 CRC64;

Query Match 90.5%; Score 19; DB 10; Length 152;
 Best Local Similarity 66.7%; Pred. No. 8.3e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
 ||| |
 Db 138 EAGTTS 143

RESULT 10
 Q9BS09 ID Q9BS09 PRELIMINARY; PRT; 161 AA.
 AC Q9BS09;
 DT 01-JUN-2001 (TEMBLrel. 17, Created)
 DT 01-JUN-2001 (TEMBLrel. 17, Last sequence update)
 DT 01-DEC-2001 (TEMBLrel. 19, Last annotation update)
 DE HYPOTHETICAL 17.1 KDA PROTEIN.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE-COLON ADENOCARCINOMA;
 RA Strausberg R.;
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; BC005805; AA05805.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 161 AA; 17058 MW; E4098AB1F0A5D706 CRC64;

Query Match 90.5%; Score 19; DB 4; Length 161;
 Best Local Similarity 66.7%; Pred. No. 8.8e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
 ||| |
 Db 43 EAGSSS 48

RESULT 11
 Q9NW11 ID Q9NW11 PRELIMINARY; PRT; 161 AA.
 AC Q9NW11;
 DT 01-OCT-2000 (TEMBLrel. 15, Created)
 DT 01-OCT-2000 (TEMBLrel. 15, Last sequence update)
 DT 01-OCT-2000 (TEMBLrel. 15, Last annotation update)
 DE CDNA FLJ20847 FIS, CLONE ADKA01746.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE-ADIPOSE TISSUE;
 RA Tanigami A., Fujiwara T., Ono T., Yamada K., Fujii Y., Ozaki K.,
 RA Hirao M., Ohmori Y., Ota T., Suzuki Y., Obayashi M., Nishi T.,
 RA Shibahara T., Tanaka T., Nakamura Y., Isogai T., Sugano S.;
 RT "NEDO human cDNA sequencing project.";
 RL Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AK000854; BAA91399.1; -.
 SQ SEQUENCE 161 AA; 17118 MW; 11098AB1EA15D71C CRC64;

Query Match 90.5%; Score 19; DB 4; Length 161;
 Best Local Similarity 66.7%; Pred. No. 8.8e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
 ||| |
 Db 43 EAGSSS 48

RESULT 12
 Q54209 ID Q54209 PRELIMINARY; PRT; 164 AA.
 AC Q54209;
 DT 01-NOV-1996 (TEMBLrel. 01, Created)
 DT 01-NOV-1996 (TEMBLrel. 01, Last sequence update)
 DT 01-DEC-2001 (TEMBLrel. 19, Last annotation update)
 DE FABD, FABH, FABC, FABB, AND ORF5 GENES.
 OS Streptomyces glaucescens.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Streptomycetaceae; Streptomyces.
 OX NCBI_TaxID=1907;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=GLA.0;
 RX MEDLINE=95352622; PubMed=7626609;
 RA Summers R.G., Ali A., Shen B., Wessel W.A., Hutchinson C.R.;
 RT "Malonyl-coenzyme A:acyl carrier protein acyltransferase of
 RT Streptomyces glaucescens: a possible link between fatty acid and
 RT polyketide biosynthesis.";
 RL Biochemistry 34:9389-9402(1995).
 DR EMBL; LA3074; AAA99450.1; -.
 SQ SEQUENCE 164 AA; 18203 MW; CB0ECF031044BB09 CRC64;

Query Match 90.5%; Score 19; DB 2; Length 164;
 Best Local Similarity 66.7%; Pred. No. 9e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxx 6
 ||| |
 Db 55 EAGTAS 60

RESULT 13
 Q9WZ11 ID Q9WZ11 PRELIMINARY; PRT; 170 AA.
 AC Q9WZ11;
 DT 01-OCT-2000 (TEMBLrel. 15, Created)
 DT 01-OCT-2000 (TEMBLrel. 15, Last sequence update)
 DT 01-JUN-2001 (TEMBLrel. 17, Last annotation update)
 DE 5-HT1A (FRAGMENT).
 OS Oryctolagus cuniculus (Rabbit).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
 OX NCBI_TaxID=9986;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=RETINA;
 RA Pootanakit K., Hunter D.D., Brunken W.J.;
 RT "5-HT1A and 5-HT7 Receptor Expression in the Mammalian Retina.";
 RL Brain Res. 0:0-0(2000); -.
 DR EMBL; AF269231; AAF76184.1; -.
 DR InterPro; IPR000276; GPCR_Rhodpsn.
 DR Pfam; PF00001; 7tm_1; 1.
 FT NON_TER 1
 FT NON_TER 170 170
 SQ SEQUENCE 170 AA; 18518 MW; 42A5B4CF917B3250 CRC64;

Query Match 90.5%; Score 19; DB 6; Length 170;
 Best Local Similarity 66.7%; Pred. No. 9.3e+02;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```
QY 1 eagxxs 6
   ||| |
Db 142 EGAAS 147

RESULT 14
Q94LT3 PRELIMINARY; PRT; 192 AA.
AC Q94LT3;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE HYPOTHETICAL 19.8 KDA PROTEIN.
OS Oryza sativa (Rice).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzeae; Oryza.
OX NCBI_TaxID=4530;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. NIPPONBARE;
RA Buell C.R., Yuan Q., Ouyang S., Moffat K.S., Hill J.N., Gansberger K.,
RA Brenner M., Burgess S., Hance M., Shvartsbeyn M., Tsitrin T.,
RA Riggs F., Hsiao J., Zismann V., Blunt S., Pal G., VanAken S.E.,
RA Utterback T.R., Feldblyum T.V., Quackenbush J., Salzberg S.L.,
RA White O., Fraser C.M.;
RT "Oryza sativa chromosome 10 BAC OSJNBb0011A08 genomic sequence.";
RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC034258; AAK54288.1; -.
KW Hypothetical protein.
SQ SEQUENCE 192 AA; 19819 MW; 4CE8C88AE83DF374 CRC64;

Query Match 90.5%; Score 19; DB 10; Length 192;
Best Local Similarity 66.7%; Pred. No. 1.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
   ||| |
Db 40 EGAAS 45

RESULT 15
Q18144 PRELIMINARY; PRT; 200 AA.
AC Q18144;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE HYPOTHETICAL 22.0 KDA PROTEIN.
GN C25A8.2.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Latreille P., Stellyes L.;
RT "The sequence of C. elegans cosmid C25A8.";
RL Science 282:2012-2018(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Latreille P., Stellyes L.;
RT "The sequence of C. elegans cosmid C25A8.";
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Waterston R.;

"Direct Submission.";
RT Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; U61958; AAB03180.1; -.
KW Hypothetical protein.
SQ SEQUENCE 200 AA; 22012 MW; 66A23EDA709C66B2 CRC64;

Query Match 90.5%; Score 19; DB 5; Length 200;
Best Local Similarity 66.7%; Pred. No. 1.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
   ||| |
Db 109 EGAAS 114

RESULT 16
O62323 PRELIMINARY; PRT; 201 AA.
AC O62323;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE R02D5.7 PROTEIN.
GN R02D5.7.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RA Matthews L.;
RL Submitted (AUG-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=99069613; PubMed=9851916;
RA none;
RT "Genome sequence of the nematode C. elegans: A platform for
RT investigating biology.";
RL Science 282:2012-2018(1998).
DR EMBL; Z78015; CAB01436.1; -.
SQ SEQUENCE 201 AA; 22266 MW; EC0423A8D7DDE4FE CRC64;

Query Match 90.5%; Score 19; DB 5; Length 201;
Best Local Similarity 66.7%; Pred. No. 1.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
   ||| |
Db 110 EGAAS 115

RESULT 17
O53374 PRELIMINARY; PRT; 204 AA.
AC O53374;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE HYPOTHETICAL 21.6 KDA PROTEIN.
GN MOA3 OR RV3322C OR MTV016.22C.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RX MEDLINE=98295987; PubMed=9634230;
RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekala F.,
RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
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RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
RA Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
RT "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence.";
RL Nature 393:537-544(1998).
DR EMBL; AL021841; CAAL17094.1; -.
DR TubercuList; RV3322c; -.
DR InterPro; IPR000051; SAM_bind.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 204 AA; 21614 MW; 13C5CB74C9C4B07F CRC64;

Query Match 90.5%; Score 19; DB 16; Length 204;
Best Local Similarity 66.7%; Pred. No. 1.1e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 153 EAGTAS 158

RESULT 18
ID Q92D00 PRELIMINARY; PRT; 212 AA.
AC Q92D00;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE LIN1021 PROTEIN.
GN LIN1021.
OS Listeria innocua.
OC Bacteria; Firmicutes; Bacillus/clostridium group;
OC Bacillus/Staphylococcus group; Listeria.
OX NCBI_TaxID=1642;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CLIP 11262 / SEROVAR 6A;
RX PubMed=11679669;
RA Glaser P., Frangeul L., Buchrieser C., Rusniok C., Amend A.,
RA Baquero F., Berche P., Bloeker H., Brandt P., Chakraborty T.,
RA Charbit A., Chetouani F., Couve E., de Daruvar A., Dehoux P.,
RA Domann E., Dominguez-Bernal G., Duchaud E., Durant L., Dussurget O.,
RA Entian K.-D., Fsihi H., Garcia-del Portillo F., Garrido P.,
RA Gautier L., Goebel W., Gomez-Lopez N., Hain T., Hauf J., Jackson D.,
RA Jones L.-M., Kaerst U., Kreft J., Kuhn M., Kunst F., Kurapkak G.,
RA Maqueno E., Maitournam A., Mata Vicente J., Ng E., Nedjari H.,
RA Nordstiek G., Novella S., de Pablo B., Perez-Diaz J.-C., Purcell R.,
RA Remmel B., Rose M., Schlueter T., Simoes N., Tierrez A.,
RA Vazquez-Boland J.-A., Voss H., Wehlund J., Cossart P.;
RT "Comparative genomics of Listeria species.";
RL Science 294:849-852(2001).
DR EMBL; AL596167; CAC96252.1; -.
DR ListiList; LIN1021; -.
KW Complete proteome.
SQ SEQUENCE 212 AA; 23672 MW; CB73DB2965A08F99 CRC64;

Query Match 90.5%; Score 19; DB 16; Length 212;
Best Local Similarity 66.7%; Pred. No. 1.2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 95 EAGASS 100

RESULT 19
ID Q91306 PRELIMINARY; PRT; 231 AA.
AC Q91306;

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DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-2001 (TrEMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE MYOSIN VI (FRAGMENT).
OS Rana catesbeiana (Bull. frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Neobatrachia; Ranioidea; Rana.
OX NCBI_TaxID=8400;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=SACCULE;
RA Solc C.F., Derfler B.H., Duyk G.M., Corey D.P.;
RT "Molecular cloning of myosins from the bullfrog sacculus macula: A
RT candidate for the hair-cell adaptation motor.";
RL Aud. Neurosci. 1:63-75(1994).
DR EMBL; U14380; AAA65089.1; -.
DR HSP; P08799; IMND.
DR InterPro; IPR001609; myosin_head.
DR Pfam; PF00063; myosin_head; 2.
DR ProDom; PD000355; myosin_head; 1.
FT NON_TER 1
FT NON_TER 231
SQ SEQUENCE 231 AA; 25693 MW; D3FFF5C343E6FAC8 CRC64;

Query Match 90.5%; Score 19; DB 13; Length 231;
Best Local Similarity 66.7%; Pred. No. 1.3e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 124 EAGSTS 129

*RESULT 20
Q9HST1
ID Q9HST1 PRELIMINARY; PRT; 245 AA.
AC Q9HST1;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE L-ISOASPARTYL PROTEIN CARBOXYL METHYLTRANSFERASE.
GN PIMT1 OR VNG0089G.
OS Halobacterium sp. (strain NRC-1).
OC Archaea; Euryarchaeota; Halobacteriales; Halobacteriaceae;
OC Halobacterium.
OX NCBI_TaxID=64091;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=2050483; PubMed=11016950;
RA Ng W.V., Kennedy S.P., Mahairas G.G., Berquist B., Pan M.,
RA Shukla H.D., Lasky S.R., Baliga N.S., Thorsson V., Sbrogna J.,
RA Swartzell S., Weir D., Hall J., Dahl T.A., Welti R., Goo Y.A.,
RA Leithausen B., Keller K., Cruz R., Danson M.J., Hough D.W.,
RA Maddocks D.G., Jablonski P.E., Krebs M.P., Angevine C.M., Dale H.,
RA Isenbarger T.A., Peck K.F., Pohlschroder M., Spudis J.L., Jung K.-H.,
RA Alam M., Freitas T., Hou S., Daniels C.J., Dennis P.P., Omer A.D.,
RA Ehardt H., Lowe T.M., Liang P., Riley M., Hood L., DasSarma S.;
RT "Genome sequence of Halobacterium species NRC-1.";
RL Proc. Natl. Acad. Sci. U.S.A. 97:12176-12181(2000).
DR EMBL; AE004377; AAC18721.1; -.
DR InterPro; IPR000682; PCMT.
DR InterPro; IPR000051; SAM_bind.
DR Pfam; PF01135; PCMT; 1.
KW Transferase; Methyltransferase; Complete proteome.
SQ SEQUENCE 245 AA; 26216 MW; A26FBBBCFAA5DB78 CRC64;

Query Match 90.5%; Score 19; DB 17; Length 245;
Best Local Similarity 66.7%; Pred. No. 1.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |

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Db 151 EAGAA 156
|||||
RESULT 21
Q91307
ID Q91307 PRELIMINARY; PRT; 254 AA.
AC Q91307;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE MYOSIN VI (FRAGMENT);
OS Rana catesbeiana (Bull frog);
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Ranidae; Rana.
OX NCBI_TaxID=8400;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=SACCULE;
RA Solc C.F., Derfler B.H., Duyk G.M., Corey D.P.;
RT "Molecular cloning of myosins from the bullfrog sacculus macula: A
candidate for the hair-cell adaptation motor.";
RL Aud. Neurosci. 1:63-75(1994).
DR EMBL; GI4381; AAA65090.1; -.
DR HSSP; PI0587; IBR2.
DR InterPro: IPR001609; myosin_head.
DR Pfam: PF00063; myosin_head; 3.
DR ProDom: PD000355; myosin_head; 1.
FT NON_TER 1
FT NON_TER 254
FT NON_TER 254
SQ SEQUENCE 254 AA; 29039 MW; DB839586BD6DE93 CRC64;

Query Match 90.5%; Score 19; DB 13; Length 254;
Best Local Similarity 66.7%; Pred. No. 1.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
|||
Db 169 EAGST 174

RESULT 22
Q9XF64
ID Q9XF64 PRELIMINARY; PRT; 257 AA.
AC Q9XF64;
DT 01-NOV-1999 (TrEMBLrel. 12, Created)
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE RING-H2 ZINC FINGER PROTEIN ATL5.
GN ATL5.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=99408259; PubMed=10480382;
RA Salinas-Mondragon R.E., Garciduenas-Pina C., Guzman P.;
RT "Early elicitor induction in members of a novel multigene family
coding for highly related RING-H2 proteins in Arabidopsis thaliana.";
RL Plant Mol. Biol. 40:579-590(1999).
CC -1- SIMILARITY: CONTAINS 1 RING-TYPE ZINC FINGER.
DR EMBL; AF132015; AAD33583.1; -.
DR InterPro: IPR001841; Znf_ring.
DR Pfam; PF00097; zf-C3HC4; 1.
DR SMART; SM00184; RING; 1.
KW Zinc-finger.
SQ SEQUENCE 257 AA; 28608 MW; 07BCE68CEC928C96 CRC64;

Query Match 90.5%; Score 19; DB 10; Length 257;
Best Local Similarity 66.7%; Pred. No. 1.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
|||
Db 184 EAGSS 189

RESULT 23
Q9LZJ6
ID Q9LZJ6 PRELIMINARY; PRT; 257 AA.
AC Q9LZJ6;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE RING-H2 ZINC FINGER PROTEIN ATL5.
GN P26K9_120.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA Blöcker H., Mewes H.W., Rudd S., Lemcke K., Mayer K.F.X., Quetier F.,
RA Salanoubat M.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA EU Arabidopsis sequencing project;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: CONTAINS 1 RING-TYPE ZINC FINGER.
DR EMBL; AL162651; CAB83119.1; -.
DR InterPro: IPR001841; Znf_ring.
DR Pfam; PF00097; zf-C3HC4; 1.
DR SMART; SM00184; RING; 1.
KW Zinc-finger.
SQ SEQUENCE 257 AA; 28592 MW; B6B7595DFF528431 CRC64;

Query Match 90.5%; Score 19; DB 10; Length 257;
Best Local Similarity 66.7%; Pred. No. 1.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
|||
Db 184 EAGSS 189

RESULT 24
Q99RZ3
ID Q99RZ3 PRELIMINARY; PRT; 260 AA.
AC Q99RZ3;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE MODA PROTEIN.
GN MODA OR SA2074.
OS Staphylococcus aureus (strain N315).
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Staphylococcus.
OX NCBI_TaxID=158879;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21311952; PubMed=11418146;
RA Kuroda M., Ohta T., Uchiyama I., Baba T., Yuzawa H., Kobayashi I.,
RA Cui L., Oguchi A., Aoki K.I., Nagai Y., Lian J., Ito T., Kanamori M.,
RA Matsumaru H., Maruyama A., Murakami H., Hosoyama A., Mizutani-Ui Y.,
RA Takahashi N.K., Sawano T., Inoue R.I., Kaito C., Sekimizu K.,
RA Hirakawa H., Kuhara S., Goto S., Yabuzaki J., Kanehisa M.,
RA Yamashita A., Oshima K., Furuya K., Yoshino C., Shiba T., Hattori M.,
RA Ogasawara N., Hayashi H., Hiramatsu K.;
RT "Whole genome sequencing of methicillin-resistant Staphylococcus
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RT aureus.";
RL Lancet 357:1225-1240(2001).
DR EMBL; AP003136; BAB43371.1; -.
DR HSSP; P37329; 1WOD.
KW Complete proteome.
SQ SEQUENCE 260 AA; 7A5D4A01A4482C4D CRC64;

Query Match          90.5%; Score 19; DB 16; Length 260;
Best Local Similarity 66.7%; Pred. No. 1.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
Db 227 EAGATS 232

RESULT 25
Q92IN7 PRELIMINARY; PRT; 261 AA.
AC Q92IN7;
DT 01-MAY-1999 (TREMBLrel. 10, Created)
DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE MODA.
GN MODA.
OS Staphylococcus carnosus.
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Staphylococcus.
OX NCBI_TaxID=1281;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-TW300;
RX MEDLINE=98340502; PubMed=9675851;
RA Neubauer H., Pantel I., Gotz F.;
RT "Characterization of moeb-part of the molybdenum cofactor
RL biosynthesis gene cluster in Staphylococcus carnosus.";
RL FEMS Microbiol. Lett. 164:55-62(1998).
DR EMBL; AF109295; AAC83133.1; -.
DR HSSP; P37329; 1WOD.
SQ SEQUENCE 261 AA; 29203 MW; 126A2D314BBAFB13 CRC64;

Query Match          90.5%; Score 19; DB 2; Length 261;
Best Local Similarity 66.7%; Pred. No. 1.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
Db 228 EAGATS 233

RESULT 26
Q92NH7 PRELIMINARY; PRT; 261 AA.
AC Q92NH7;
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE PUTATIVE TRANSCRIPTION REGULATOR PROTEIN.
GN SMC01615.
OS Rhizobium meliloti (Sinorhizobium meliloti).
OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
OC Rhizobiaceae; Sinorhizobium.
OX NCBI_TaxID=382;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=1021;
RX MEDLINE=21368234; PubMed=11474104;
RA Galibert F., Finan T.M., Long S.R., Puehler A., Abola P., Ampe F.,
RA Barloy-Hubler F., Barnett M.J., Becker A., Boistard P., Bothe G.,
RA Boutry M., Bowser L., Buhrmester J., Cadieu E., Capela D., Chain P.,
RA Cowie A., Davis R.W., Dreano S., Federspiel N.A., Fisher R.F.,

RA Gloux S., Godrie T., Goffeau A., Golding B., Gouzy J., Gurjal M.,
RA Hernandez-Lucas I., Hong A., Huizar L., Hyman R.W., Jones T., Kahn D.,
RA Kahn M.L., Kalman S., Keating D.H., Kiss E., Komp C., Lelaure V.,
RA Masuy D., Palm C., Peck M.C., Pohl T.M., Portetelle D., Purnelle B.,
RA Ramsberger U., Surzycki R., Thebault P., Vandebol M.,
RA Vorhoefer F.J., Weidner S., Wells D.H., Wong K., Yeh K.-C., Batut J.;
RT "The composite genome of the legume symbiont Sinorhizobium meliloti.";
RL Science 293:668-672(2001).
DR EMBL; AL591790; CAC46806.1; -.
KW Complete proteome.
SQ SEQUENCE 261 AA; 28115 MW; 031D03708E1084CB CRC64;

Query Match          90.5%; Score 19; DB 16; Length 261;
Best Local Similarity 66.7%; Pred. No. 1.4e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
Db 16 EAGTAS 21

RESULT 27
Q88190 PRELIMINARY; PRT; 266 AA.
ID Q88190
AC Q88190;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE COAT PROTEIN.
OS Soybean mosaic virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Potyviridae;
OX NCBI_TaxID=12222;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-SA;
RA Chu R., Leng X., Bao Y., Pu Z., Pan N., Chen Z.;
RT "Amplification of soybean mosaic virus coat protein gene by polymerase
RL chain reaction and its sequence analysis.";
RL Acta Bot. Sin. 34:523-528(1992).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-SA;
RA Xu L.;
RL Submitted (APR-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; U25673; AAA70095.1; -.
DR InterPro; IPR001592; Poty_coat.
DR Pfam; PF00767; Poty_coat; 1.
SQ SEQUENCE 266 AA; 30084 MW; 4E08AFE7D434307F CRC64;

Query Match          90.5%; Score 19; DB 12; Length 266;
Best Local Similarity 66.7%; Pred. No. 1.5e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
Db 25 EAGTSS 30

RESULT 28
Q88196 PRELIMINARY; PRT; 267 AA.
ID Q88196
AC Q88196;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE COAT PROTEIN (FRAGMENT).
OS Soybean mosaic virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Potyviridae;
OX NCBI_TaxID=12222;

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RN  [1]
RP  SEQUENCE FROM N.A.
RC  STRAIN=CHINESE;
RA  Chu R.;
RT  *CDNA sequence of the gene encoding coat protein of SMV. ";
RL  Nucleic Acids Res. 0:0-0(0).
DR  EMBL; X63771; CAA45307.1; -.
DR  InterPro; IPR001592; Poty_coat.
DR  Pfam; PF00767; Poty_coat; 1.
FT  NON_TER
FT  1
SQ  SEQUENCE 267 AA; 30104 MW; 220E42F2595BE059 CRC64;

Query Match          90.5%; Score 19; DB 12; Length 267;
Best Local Similarity 66.7%; Pred. No. 1.5e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  1 eaqxgs 6
DB  26 EAGTSS 31

RESULT 29
Q9DCE4 PRELIMINARY; PRT; 269 AA.
AC Q9DCE4;
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DE 0610039P13RIK PROTEIN.
GN Mus musculus (Mouse).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
[1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=KIDNEY;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
RA Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli K., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
DR EMBL; AK002854; BAB22409.1; -.
DR MGD; MGI:1921346; 0610039P13RIK.
DR InterPro; IPR000636; Cation_chan_non_lig.
SQ SEQUENCE 269 AA; 31242 MW; B549CB553DEB6568 CRC64;

Query Match          90.5%; Score 19; DB 11; Length 269;
Best Local Similarity 66.7%; Pred. No. 1.5e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY  1 eaqxgs 6
DB  71 EAGSAS 76

RESULT 30
Q9VGK4 PRELIMINARY; PRT; 270 AA.
AC Q9VGK4;
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE CG14714 PROTEIN.
GN CG14714.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
[1]
RP SEQUENCE FROM N.A.
RC STRAIN=BERKELEY;
RX MEDLINE=20196006; PubMed=10731132;
RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Ananides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
RA Abril J.F., Aghayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
RA Balleg R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brotter P.,
RA Burtis K.C., Busam D.A., Butler H., Cadiu E., Center A., Chandra I.,
RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K.J., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
RA Foster C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Helman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwac C.,
RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattel B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wassarman D.A., Weinstein G.M., Weissbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2185-2195(2000).
DR EMBL; AE003692; AAF54673.1; -.
DR HSSP; P28827; IRPM.
DR FlyBase; FBgn0037929; CG14714.
DR InterPro; IPR000387; TYR_phosphatase.
DR InterPro; IPR000242; Tyr_prot_phptase.
DR Pfam; PF00102; Y-phosphatase; 1.
DR PRINTS; PR00700; PRTYPPHTASE.
DR SMART; SM00194; PTPC; 1.
DR PROSITE; PS00383; TYR_PHOSPHATASE_1; 1.
DR PROSITE; PS50056; TYR_PHOSPHATASE_2; 1.
DR PROSITE; PS50055; TYR_PHOSPHATASE_PTP; 1.
KW Hydrolase.
SQ SEQUENCE 270 AA; 30259 MW; 5C136F3135CAD001 CRC64;

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Query Match

90.5%; Score 19; DB 5; Length 270;

Best Local Similarity 66.7%; Pred. No. 1.5e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 169 EAGSTS 174

RESULT 31

ID Q9A1T5 PRELIMINARY; PRT; 285 AA.
AC Q9A1T5;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE ICMF-LIKE PROTEIN (FRAGMENT).
OS Vibrio cholerae.
OC Bacteria; Proteobacteria; gamma subdivision; Vibrionaceae; Vibrio.
OX NCBI_TaxID=666;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=569B;
RX MEDLINE=20434574; PubMed=10981695;
RA Das S., Chakraborty A., Banerjee R., Roychoudhury S., Chaudhuri K.;
RT "Comparison of global transcription responses allows identification of
RT Vibrio cholerae genes differentially expressed following infection.";
RL FEMS Microbiol. Lett. 190:87-91(2000).
DR EMBL: AF239737; AAK27321.1; -.
FT NON_TER 1 285
FT NON_TER 285
SQ SEQUENCE 285 AA; 32165 MW; BDDE7FA9021F5661 CRC64;

Query Match 90.5%; Score 19; DB 2; Length 285;
Best Local Similarity 66.7%; Pred. No. 1.6e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 210 EAGSAS 215

RESULT 32

ID O93503 PRELIMINARY; PRT; 287 AA.
AC O93503;
DT 01-NOV-1998 (TrEMBLrel. 08, Created)
DT 01-NOV-1998 (TrEMBLrel. 08, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE MYRISTOYLATED ALANINE-RICH C KINASE SUBSTRATE.
GN MARCKS.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98030614; PubMed=9361009;
RA Shi Y., Sullivan S.K., Pitterle D.M., Kennington E.A., Graff J.M.,
RA Blackshear P.J.;
RT "Mechanisms of MARCKS gene activation during Xenopus development.";
RL J. Biol. Chem. 272:29290-29300(1997).
DR EMBL: AF017299; AAC61897.1; -.
DR InterPro: IPR002101; MARCKS.
DR Pfam: PF02063; MARCKS; 1.
DR PRINTS; PR00963; MARCKS.
DR PROSITE; PS00826; MARCKS_1; 1.
DR PROSITE; PS00827; MARCKS_2; 1.
KW Kinase.
SQ SEQUENCE 287 AA; 29147 MW; 35CB7AE6090ED3C1 CRC64;

Query Match 90.5%; Score 19; DB 13; Length 287;
Best Local Similarity 66.7%; Pred. No. 1.6e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 110 EAGSTS 115

RESULT 33

ID Q9L866 PRELIMINARY; PRT; 293 AA.
AC Q9L866;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE HPOTHEICAL 32.1 KDA PROTEIN.
OS Azospirillum brasilense.
OC Bacteria; Proteobacteria; alpha subdivision; Rhodospirillaceae;
OX Azospirillum.
RN NCBI_TaxID=192;
RP SEQUENCE FROM N.A.
RA Ma L., Zhao Y., Wang J., Li J.;
RT "Sequence and function analysis of draTG genes downstream ORFs from
RT Azospirillum brasilense yu62.";
RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF216815; AA561911.1; -.
DR InterPro: IPR003310; DNA_glycosylase.
KW Hypothetical protein.
SQ SEQUENCE 293 AA; 32063 MW; 37417EA008F6BD61 CRC64;

Query Match 90.5%; Score 19; DB 2; Length 293;
Best Local Similarity 66.7%; Pred. No. 1.6e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
||| |
Db 168 EAGAAS 173

RESULT 34

ID Q91296 PRELIMINARY; PRT; 297 AA.
AC Q91296;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE MYOSIN VI (FRAGMENT).
OS Rana catesbeiana (Bull frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Neobatrachia; Ranidae; Rana.
OX NCBI_TaxID=8400;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=SACCULE;
RA Solc C.F., Derfler B.H., Duyk G.M., Corey D.P.;
RT "Molecular cloning of myosins from the bullfrog saccular macula: A
RT candidate for the hair-cell adaptation motor.";
RL Aud. Neurosci. 1:63-75(1994).
DR EMBL: U14370; AAA65079.1; -.
DR HSPSP; P10587; 1BR2.
DR InterPro: IPR001609; myosin_head.
DR Pfam: PF00063; myosin_head; 2.
DR ProDom: PD000355; myosin_head; 1.
FT NON_TER 1 297
FT NON_TER 297
SQ SEQUENCE 297 AA; 33755 MW; 46EE6C78A8ED530D CRC64;

Query Match 90.5%; Score 19; DB 13; Length 297;
Best Local Similarity 66.7%; Pred. No. 1.7e+03;

Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 eagxxs 6
   ||| |
Db 169 EAGSTS 174

RESULT 35
Q9PCJ6 PRELIMINARY; PRT; 302 AA.
AC Q9PCJ6;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DE 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE HYPOTHETICAL PROTEIN XF1783.
GN XF1783.
OS Xylella fastidiosa.
OC Bacteria; Proteobacteria; gamma subdivision; Xanthomonas group;
OC Xylella.
OX NCBI_TaxID=2371;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=9A5C;
RX MEDLINE=20365717; PubMed=10910347;
RA Simpson A.J.G., Reinach F.C., Arruda P., Abreu F.A., Acencio M.,
RA Alvarenga R., Alves L.M.C., Araya J.E., Baia G.S., Baptista C.S.,
RA Barros M.H., Bonaccorsi E.D., Bordin S., Bove J.M., Briones M.R.S.,
RA Bueno M.R.P., Camargo A.A., Canargo L.E.A., Carraro D.M., Carrer H.,
RA Colauto N.B., Colombo C., Costa F.F., Costa M.C.R., Costa-Neto C.M.,
RA Coutinho L.L., Cristofani M., Dias-Neto E., Docena C., El-Dorri H.,
RA Facincani A.P., Ferreira A.J.S., Ferreira V.C.A., Ferro J.A.,
RA Fraga J.S., Franca S.C., Franco M.C., Frohme M., Furlan L.R.,
RA Garnier M., Goldman G.H., Goldman M.H.S., Gomes S.L., Gruber A.,
RA Ho P.L., Hobeisel J.D., Junqueira M.L., Kemper E.L., Kitajima J.P.,
RA Krieger J.E., Kuramae E.F., Laigret F., Lambais M.R., Leite L.C.C.,
RA Lemos E.G.M., Lemos M.V.F., Lopes S.A., Lopes C.R., Machado J.A.,
RA Machado M.A., Martins E.A.L., Martins E.M.F., Matsukuma A.Y.,
RA Marques M.V., Miracca E.C., Miyaki C.Y., Monteiro-Vitorello C.B.,
RA Menck C.F.M., Miracca E.C., Miyaki C.Y., Monteiro-Vitorello C.B.,
RA Moon D.H., Nagai M.A., Nascimento A.L.T.O., Netto L.E.S.,
RA Nhani A. Jr., Nobrega F.G., Nunes L.R., Oliveira M.A.,
RA de Oliveira M.C., de Oliveira R.C., Palmieri D.A., Paris A.,
RA Peixoto B.R., Pereira G.A.G., Pereira H.A. Jr., Pesquero J.B.,
RA Quaggio R.B., Roberto P.G., Rodrigues V., de Rosa A.J.M.,
RA de Rosa V.E. Jr., de Sa R.G., Santelli R.V., Sawasaki H.E.,
RA da Silva A.C.R., da Silva A.M., da Silva F.R., Silva W.A. Jr.,
RA da Silveira J.F., Silvestri M.L.Z., Siqueira W.J., de Souza A.A.,
RA de Souza A.P., Terenzi M.F., Truffi D., Tsai S.M., Tsubako M.H.,
RA Vallada H., Van Sluys M.A., Verjovski-Almeida S., Vettore A.L.,
RA Zago M.A., Zatz M., Meidanis J., Setubal J.C.;
RT "The genome sequence of the plant pathogen Xylella fastidiosa."
RL Nature 406:151-159(2000).
DR EMBL: AE004000; AAF84591.1;
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 302 AA; 32047 MW; 758CC61DE4BE3590 CRC64;
```

Query Match 90.5%; Score 19; DB 16; Length 302;
Best Local Similarity 66.7%; Pred. No. 1.7e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 eagxxs 6
   ||| |
Db 106 EAGTAS 111
```

```
RESULT 36
Q9FJG2 PRELIMINARY; PRT; 307 AA.
AC Q9FJG2;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-MAR-2001 (TrEMBLrel. 16, Last annotation update)
```

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DE GB|AAD26962.1.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eucosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=COLUMBIA;
RX MEDLINE=99087489; PubMed=9872454;
RA Nakamura Y., Sato S., Asamizu E., Kaneko T., Kotani H., Miyajima N.,
RA Tabata S.;
RT "Structural analysis of Arabidopsis thaliana chromosome 5. VII.
RT Sequence features of the regions of 1,013,767 bp covered by sixteen
RT physically assigned P1 and TAC clones.";
RL DNA Res. 5:297-308(1998).
DR EMBL: AB015473; BAB08399.1; -.
SQ SEQUENCE 307 AA; 35727 MW; 23CDB6C127CE90D1 CRC64;
```

Query Match 90.5%; Score 19; DB 10; Length 307;
Best Local Similarity 66.7%; Pred. No. 1.7e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 eagxxs 6
   ||| |
Db 207 EAGTSS 212
```

```
RESULT 37
Q90888 PRELIMINARY; PRT; 311 AA.
ID Q90888;
AC Q90888;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DE 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE MAFB.
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianinae;
OC Gallus.
OX NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=95021288; PubMed=7935473;
RA Kataoka K., Fujiwara K.T., Noda M., Nishizawa M.;
RT "WafB, a new Maf family transcription activator that can associate with
RT Maf and Fos but not with Jun.";
RL Mol. Cell. Biol. 14:7581-7591(1994).
DR EMBL: D28600; BAA05938.1; -.
DR InterPro: IPR001871; bZIP.
DR Pfam: PF03131; bZIP_Maf; 1.
DR SMART: SM00338; BRLZ; 1.
SQ SEQUENCE 311 AA; 35467 MW; DDAE7F698B7D3ABA CRC64;
```

Query Match 90.5%; Score 19; DB 13; Length 311;
Best Local Similarity 66.7%; Pred. No. 1.7e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 eagxxs 6
   ||| |
Db 296 EAGSTS 301
```

```
RESULT 38
Q90370 PRELIMINARY; PRT; 311 AA.
ID Q90370;
AC Q90370;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE MAFB PROTEIN.
```

GN MAFB.
 OS Coturnix coturnix japonica (Japanese quail).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
 OC Coturnix.
 OX NCBI_TaxID=93934;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=96180718; PubMed=8620536;
 RA Stewek M.H., Tekotte H., Frampton J., Graf T.;
 RT "MafB is an interaction partner and repressor of Ets-1 that inhibits
 erythroid differentiation.";
 RL Cell 85:49-60(1996).
 DR EMBL; X96511; CAA65360.1; -.
 DR InterPro; IPR001871; bZIP.
 DR Pfam; PF03131; bZIP_Maf; 1.
 DR SMART; SM00338; BRUZ; 1.
 SQ SEQUENCE 311 AA; 35476 MW; 7D1F3FA05D5CD683 CRC64;

Query Match 90.5%; Score 19; DB 13; Length 311;
 Best Local Similarity 66.7%; Pred. No. 1.7e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 296 EAGSTS 301

RESULT 39

ID Q9PUA6 PRELIMINARY; PRT; 313 AA.
 AC Q9PUA6;
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE bZIP TRANSCRIPTION FACTOR MAFB.
 GN MAFB.
 OS Xenopus laevis (African clawed frog).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
 OC Xenopodinae; Xenopus.
 OX NCBI_TaxID=8355;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21152895; PubMed=11231066;
 RA Ishibashi S., Yasuda K.;
 RT "Distinct roles of maf genes during Xenopus lens development.";
 RL Mech. Dev. 101:155-166(2001).
 DR EMBL; AF202058; AAF08316.1; -.
 DR InterPro; IPR001871; bZIP.
 DR Pfam; PF03131; bZIP_Maf; 1.
 DR SMART; SM00338; BRUZ; 1.
 SQ SEQUENCE 313 AA; 35714 MW; 8E697A00A928BF95 CRC64;

Query Match 90.5%; Score 19; DB 13; Length 313;
 Best Local Similarity 66.7%; Pred. No. 1.7e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 298 EAGSTS 303

RESULT 40

ID Q9Y5Q3 PRELIMINARY; PRT; 323 AA.
 AC Q9Y5Q3;
 DT 01-NOV-1999 (TrEMBLrel. 12, Created)
 DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE MAFB/KREISLER BASIC REGION/LEUCINE ZIPPER TRANSCRIPTION FACTOR.

GN MAFB.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX TISSUE=BONE MARROW;
 RX MEDLINE=99375320; PubMed=10444328;
 RA Wang P.W., Eisenbart J.D., Cordes S.P., Barsh G.S., Stoffel M.,
 RA Le Beau M.M.;
 RT "Human KRML (MAFB): cDNA cloning, genomic structure, and evaluation as
 a candidate tumor suppressor gene in myeloid leukemias.";
 RL Genomics 59:275-281(1999).
 DR EMBL; AF134157; AAD30106.1; -.
 DR InterPro; IPR001871; bZIP.
 DR Pfam; PF03131; bZIP_Maf; 1.
 DR SMART; SM00338; BRUZ; 1.
 SQ SEQUENCE 323 AA; 35829 MW; AB4DC23408E36E55 CRC64;

Query Match 90.5%; Score 19; DB 4; Length 323;
 Best Local Similarity 66.7%; Pred. No. 1.8e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 308 EAGSTS 313

RESULT 41

ID Q9H1F1 PRELIMINARY; PRT; 323 AA.
 AC Q9H1F1;
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)
 DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE DJ64411.1 (KREISLER (MOUSE) MAF-RELATED LEUCINE ZIPPER HOMOLOGY).
 GN KRML.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Ramsay H.;
 RL Submitted (MAR-2001) to the EMBL/GenBank/DBDJ databases.
 DR EMBL; AL035665; CAB75863.1; -.
 DR InterPro; IPR001871; bZIP.
 DR SMART; SM00338; BRUZ; 1.
 SQ SEQUENCE 323 AA; 35792 MW; A0F3C09F8936CB16 CRC64;

Query Match 90.5%; Score 19; DB 4; Length 323;
 Best Local Similarity 66.7%; Pred. No. 1.8e+03;
 Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxs 6
 ||| |
 Db 308 EAGSTS 313

RESULT 42

ID Q98TS3 PRELIMINARY; PRT; 333 AA.
 AC Q98TS3;
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE HSF70 BINDING PROTEIN.
 OS Brachydanio rerio (zebrafish) (zebra danio).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Osteichthyes; Ostariophysi;

OC Cypriniformes: Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RA Guerrier V., Raynes D.A.;
RT "Hsp70 binding protein from zebra fish (HspBPF).";
RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY024336; AAG61257.1; -
DR InterPro: IPR000225; Armadillo.
DR Pfam: PF00514; Armadillo_seg; 2.
DR SMART: SM00185; ARM; 2.
SQ SEQUENCE 333 AA; 37269 MW; E7C5ABD12F41D23E CRC64;

Query Match 90.5%; Score 19; DB 13; Length 333;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 23 EAGSAS 28

RESULT 43
Q9ATS6 PRELIMINARY; PRT; 346 AA.
ID Q9ATS6;
AC Q9ATS6;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE TEOSINTE BRANCHED PROTEIN (FRAGMENT).
GN TBL.
OS Arundinella hirta.
OC Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC clade;
OC Panicoideae; Arundinelleae; Arundinella.
OX NCBI_TaxID=79825;

SEQUENCE FROM N.A.
RN [1]
RX MEDLINE=21165336; PubMed=11264415;
RA Lukens L., Doebley J.;
RT "Molecular evolution of the teosinte branched gene among maize and related grasses.";
RL Mol. Biol. Evol. 18:627-638(2001).
DR EMBL; AF322131; AAK37493.1; -
FT NON_TER 1
FT NON_TER 346
SQ SEQUENCE 346 AA; 36902 MW; BFB5F29CD7449C89 CRC64;

Query Match 90.5%; Score 19; DB 10; Length 346;
Best Local Similarity 66.7%; Pred. No. 1.9e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 280 EAGAS 285

RESULT 44
Q98IH9 PRELIMINARY; PRT; 350 AA.
ID Q98IH9;
AC Q98IH9;
DT 01-OCT-2001 (TREMBlrel. 18, Created)
DT 01-OCT-2001 (TREMBlrel. 18, Last sequence update)
DT 01-OCT-2001 (TREMBlrel. 18, Last annotation update)
DE MLR9369 PROTEIN.
GN MLR9369.
OS Rhizobium loti (Mesorhizobium loti).
OG Plasmid pMLA.
OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
OC Phyllobacteriaceae; Mesorhizobium.
OX NCBI_TaxID=381;

RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=MAFF303099;
RA MEDLINE=21082930; PubMed=11214968;
RA Kaneko T., Nakamura Y., Sato S., Asamizu E., Kato T., Sasamoto S.,
RA Watanabe A., Idesawa K., Ishikawa A., Kawashima K., Kimura T.,
RA Kishida Y., Kiyokawa C., Kohara M., Matsumoto M., Matsumoto A.,
RA Mochizuki Y., Nakayama S., Nakazaki N., Shimo S., Sugimoto M.,
RA Takeuchi C., Yamada M., Tabata S.;
RT "Complete genome structure of the nitrogen-fixing symbiotic bacterium Mesorhizobium loti.";
RL DNA Res. 7:331-338(2000).
DR EMBL; AF003016; BAB54976.1; -
KW Plasmid; Complete proteome.
SQ SEQUENCE 350 AA; 39172 MW; BB075F9345BB9362 CRC64;

Query Match 90.5%; Score 19; DB 16; Length 350;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 108 EAGASS 113

RESULT 45
Q92WT2 PRELIMINARY; PRT; 352 AA.
ID Q92WT2;
AC Q92WT2;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-DEC-2001 (TREMBlrel. 19, Last annotation update)
DE PUTATIVE ADENYLATE CYCLASE PROTEIN.
GN SMB2025.
OS Rhizobium meliloti (Sinorhizobium meliloti).
OG Plasmid pSymb (megaplasmid 2).
OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
OC Rhizobiaceae; Sinorhizobium.
OX NCBI_TaxID=382;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=1021;
RX MEDLINE=21396508; PubMed=11481431;
RA Finan T.M., Weidner S., Wong K., Buhrmester J., Chain P.,
RA Vorhoelter F.J., Hernandez-Lucas I., Becker A., Cowie A., Gouzy J.,
RA Golding B., Puehler A.;
RT "The complete sequence of the 1.683-kb pSymb megaplasmid from the N2-fixing endosymbiont Sinorhizobium meliloti.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:9889-9894(2001).
DR EMBL; AL603642; CAC48647.1; -
KW Plasmid; Hypothetical protein; Complete proteome.
SQ SEQUENCE 352 AA; 37018 MW; F4AE6710196E06EF CRC64;

Query Match 90.5%; Score 19; DB 16; Length 352;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
||| |
Db 13 EAGTSS 18

RESULT 46
Q98UK5 PRELIMINARY; PRT; 356 AA.
ID Q98UK5;
AC Q98UK5;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-DEC-2001 (TREMBlrel. 19, Last annotation update)
DE TRANSCRIPTION FACTOR MAFB.
GN MAFB.

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OS Brachydanio rerio (Zebrafish) (Zebra danio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Ostariophysi;
OC Cypriniformes; Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21064923; PubMed=11134968;
RA Kajihara M., Kawauchi S., Kobayashi M., Ogino H., Takahashi S.,
RA Yasuda K.;
RT "Isolation, Characterization, and Expression Analysis of Zebrafish
RT Large Maf."
RL J. Biochem. 129:139-146(2001).
DR EMBL; AB006322; BAB21102.1; -.
DR InterPro; IPR001871; bZIP.
DR Pfam; PF03131; bZIP_Maf; 1.
DR SMART; SM00338; BRLZ; 1.
SQ SEQUENCE 356 AA; 40233 MW; DE4C96B62C058865 CRC64;

Query Match          90.5%; Score 19; DB 13; Length 356;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 eagxxx 6
Db 341 EAGSTS 346

RESULT 47
O73679 PRELIMINARY; PRT; 356 AA.
AC O73679;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE TRANSCRIPTION FACTOR VAL.
GN VAL OR VALENTINO.
OS Brachydanio rerio (zebrafish) (Zebra danio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Ostariophysi;
OC Cypriniformes; Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98165393; PubMed=9425134;
RA Moens C.B., Cordes S.P., Giorgianni M.W., Barsh G.S., Kimmel C.B.;
RT "Equivalence in the genetic control of hindbrain segmentation in fish
RT and mouse."
RL Development 125:381-391(1998).
DR EMBL; AF006641; AAC18821.1; -.
DR ZFIN; ZDB-GENE-980526-515; val.
DR InterPro; IPR001871; bZIP.
DR Pfam; PF03131; bZIP_Maf; 1.
DR SMART; SM00338; BRLZ; 1.
SQ SEQUENCE 356 AA; 40243 MW; 07420DB0F6CD08F1 CRC64;

Query Match          90.5%; Score 19; DB 13; Length 356;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 eagxxx 6
Db 341 EAGSTS 346

RESULT 48
O23101 PRELIMINARY; PRT; 357 AA.
ID O23101
AC O23101;
DT 01-JAN-1998 (TrEMBLrel. 05, Created)
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)

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DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE A_TM018A10.10 PROTEIN.
GN A_TM018A10.10
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. COLUMBIA;
RA Dempsey S., Harper M.;
RT "The sequence of A. thaliana TM018A10."
RL Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. COLUMBIA;
RA Washu;
RT "The A. thaliana Genome Sequencing Project."
RL Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. COLUMBIA;
RA Waterston R.;
RL Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF013294; AAB62869.1; -.
DR InterPro; IPR001810; F-box.
DR Pfam; PF00646; F-box; 1.
DR SMART; SM00256; FBOX; 1.
SQ SEQUENCE 357 AA; 40078 MW; B1683A07BF630633 CRC64;

Query Match          90.5%; Score 19; DB 10; Length 357;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 eagxxx 6
Db 91 EAGSSS 96

RESULT 49
Q9BL91 PRELIMINARY; PRT; 361 AA.
ID Q9BL91
AC Q9BL91;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE HYPOTHEtical 40.2 KDA PROTEIN.
GN Y18H1A.7.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RX MEDLINE=99069613; PubMed=9851916;
RA None;
RT "Genome sequence of the nematode C. elegans: a platform for
RT investigating biology. The C. elegans Sequencing Consortium."
RL Science 282:2012-2018(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Bemis G., Lamar B., Courtney L., Wohlmann P., Harrison M.;
RT "The sequence of C. elegans cosmid Y18H1A."
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Waterston R.;
RT "Direct Submission."

```

RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC024751; AAK21510.1; -
KW Hypothetical protein.
SQ SEQUENCE 361 AA; 40201 MW; 98F44C3F87D59625 CRC64;

Query Match 90.5%; Score 19; DB 5; Length 361;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
|||
Db 80 EAGSSS 85

RESULT 50
Q9JHQ1
ID Q9JHQ1 PRELIMINARY; PRT; 362 AA.
AC Q9JHQ1;
DT 01-OCT-2000 (TReMBLrel. 15, Created)
DT 01-OCT-2000 (TReMBLrel. 15, Last sequence update)
DT 01-DEC-2001 (TReMBLrel. 19, Last annotation update)
DE TITIN (FRAGMENT).
GN TTN.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-SKELETAL MUSCLE;
RX MEDLINE=20490785; PubMed=11034912;
RA Person V., Kostin S., Suzuki K., Labeit S., Schaper J.;
RT "Antisense oligonucleotide experiments elucidate the essential role of
RT titin in sarcomerogenesis in adult rat cardiomyocytes in long-term
RT culture.";
RL J. Cell Sci. 113:3851-3859(2000).
DR EMBL; AJ401157; CAB95001.1; -
DR HSSP; P56276; 1TLK.
DR InterPro; IPR003598; Ig_c2.
DR InterPro; IPR003600; Ig_like.
DR InterPro; IPR003006; Ig_MHC.
DR Pfam; PF00047; Ig; 3.
DR SMART; SM00408; Igc2; 2.
DR SMART; SM00410; IG_like; 1.
KW Immunoglobulin domain.
FT NON_TER 1
FT NON_TER 362
SQ SEQUENCE 362 AA; 39601 MW; E8E6CE65BB7F4ED8 CRC64;

Query Match 90.5%; Score 19; DB 11; Length 362;
Best Local Similarity 66.7%; Pred. No. 2e+03;
Matches 4; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 eagxxs 6
|||
Db 317 EAGSSS 322

Search completed: August 30, 2002, 15:11:35
Job time: 346 sec

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